



Cranio-maxillofacial

Implant Directions®

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CASE REPORT

TREATMENT OF A SEVERLY RESORBED MANDIBLE WITH ENDOSSEOUS IMPLANTS IN AN IMMEDIATE LOADING PROTOCOL

EVIDENCE REPORT

A COMPARISON OF BONE GRAFTS WITH AND WITHOUT PLATELET-RICH PLASMA IN PREPARATION FOR DENTAL IMPLANT PLACEMENT

RESEARCH IN CONTEXT

DEVELOPING THE STUDY PLAN

FULL LENGTH ARTICLE

FUNCTIONALLY RELATED CHANGES TO THE VERTICAL PERIMPLANT BONE AFTER BOI IMPLANT INSERTION IN MANDIBLE

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Contact

publishing@implantfoundation.org

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- Evidence Reports summarize the latest «Hot Topics» from relevant journals putting similar studies «side-by-side». This unique presentation of studies allows you to compare and contrast the patient populations, the treatment interventions, and the quality of the scientific methods. The «evidence-based bottom line» is presented with an overall summary statement at the beginning. Clinical notes by implantologists with special expertise on the topic complete the Evidence Report by providing their expert clinical opinion. ID is an implantology publication that provides attention to detail in balancing science with clinical opinion in such a clear, concise, and visually-friendly presentation.
- Literature Analyses provide you with an in-depth look at the research on a given topic. A «Literature Analysis» is a critical review of the literature on the epidemiology, treatment methods, and prognosis for implant-related topics or conditions. Literature Analyses are broader than «Evidence Reports» and are written to serve as a reference tool for implantologists to help them make decisions regarding how to manage patients, to assist them in evaluating needs for future research, and to use the material for future presentations.
- Critical Appraisals summarize the findings from important papers used for clinical decision making or marketing by implant companies. In addition to the summary, the study's methods and clinical conclusions are critically reviewed in an effort to challenge the implantology community into not accepting everything that is published, while fostering alternative explanations and ideas.
- Case reports give implantologists the opportunity to publish on unique patients using innovative or alternative methods for treating challenging patient conditions.
- Research in Context is a helpful «what is» section to consult if you've ever read a study and asked «what is a p-value» or any other research method question. It assists clinicians with the critical evaluation of the literature by briefly describing relevant aspects of research methods and statistical analysis that may bias results and lead to erroneous conclusions.

A case report.

Treatment of a severely resorbed mandible with endosseous implants in an immediate loading protocol

Authors:

Dr. Antonina Rusak
V. Choruzhej Str. 34a-374
BY-220123 Minsk
Belarus
Dr.Rusak@mail.ru

Dr. med.dent. S.K.A. Ihde
MNE-85386 Vrba/Tudorovici, Montenegro
Email: ihde@ihde.com

Corresponding author: Dr. Antonina Rusak

Abstract

This article reports on the implantological treatment in the atrophied lower jaw. To avoid augmentations and nevertheless equip the distal mandible with endosseous implants, we have used basal implants in such a way, that the base plates were positioned below the alveolar nerve in the region of the canines, and above the nerve distally to the area of the 2nd molar. All implants were immediately splinted and thereby loaded with a circular bridge. The aesthetic and functional outcome was satisfactory. By using this technique a secure distal bridge support is available for almost all patients. The placement of base-plates below the lower alveo-

lar nerve is a useful technique.

Keywords:

Basal implants, Diskos[®], treatment of mandibular atrophy, immediate loading, avoiding augmentations.

Introduction

The conventional dental implant treatment imposes difficulties, when it comes to treat the severely atrophied mandible. This case report demonstrates advantages of a treatment approach with basal implants

Material and Method

A 65-year-old female patient without any generalized diseases requested implant treatment in the mandible. The patient was edentulous in both jaws. The preoperative panoramic view (Fig. 1) revealed a pronounced atrophy. After evaluating all alternatives, it was decided by the patient to undergo treatment with basal implants. To overcome the problems caused by the atrophy and allow the installation of a wider load transmission areas, the base plates of both anterior implants were positioned below the alveolar nerve , .

Technique:

In local anasesthesia a wide flaps were prepared on each side of the mandible to allow surgical access from the lateral aspect. The vertical slot for the mesial basal implants was prepared. Into

the bottom part of this osteotomy the horizontal slot for the base plate was prepared to a width of 9mm and a height of 0.6 mm. Basal

implants require vestibular and lingual engagement. Therefore the necessary diameter of the plate is determined by the clinical situation.

After placing two more implants distally, the flap was closed and sutured. Healing was uneventful, no paraesthesia was reported. The definitive bridge was delivered on day three postoperatively. Fig. 2 shows the case 8 years postoperatively

Conclusion

The usage of basal implants allows treating also those patients in a single surgical approach, which need the treatment most. The procedure avoids the burdens and costs of bone augmentations and leads to immediate function and customer satisfaction. Placing the base plates below the mandibular nerve makes the treatment more practical. The width of the mandible instead of the height is utilized for stabilizing these implants. Basal implants are safe and effective treatment devices , , , . Their use should be considered when patients with atrophied jaw bones request dental implant treatment.

References

Ihde S. Principles of BOI, Springer, 2005 Heidelberg/New-York

J.M. Donsimoni, P. Bermot, D. Dohan: Les implants maxillo-faciaux à plateaux d'assise Concepts et technologies orthopédiques, réhabilitations maxillo-mandibulaires, reconstructions maxillo-faciales, réhabilitations dentaires partielles, techniques de réintervention, méta-analyse. 2e partie : réhabilitations maxillo-mandibulaires. *Implantodontie* 13 (2004) 31-41

Ihde (2008): Outcomes of immediately loaded full arch reconstructions on basal implants in the mandible: retrospective report on 115 consecutive cases during a period of up to 134 months. *CMF Impl. Dir* 2008(3) 1: 50-60

Kopp (2007): Basal implants: a safe and effective treatment option in dental implantology; *CMF Impl. Dir. Vol.2 No. 3*, p. 110-117

Scortecchi G, Misch C, Benner K (2001): Implants and restorative dentistry. Dunitz, London (Textbook publication including statistics)

Ihde S, Mutter E (2003): Versorgung von Freisituationen mit basalosseointegrierten Implantaten (BOI) bei reduziertem vertikalem Knochenangebot. *Dtsch Zahnärztl Z*, 58:94–102.

Conflict of interest statement

Prof. Dr. Stefan Ihde declares that a potential conflict of interest may exist due to his employment in the Dr. Ihde Dental group of companies (www.implant.com)

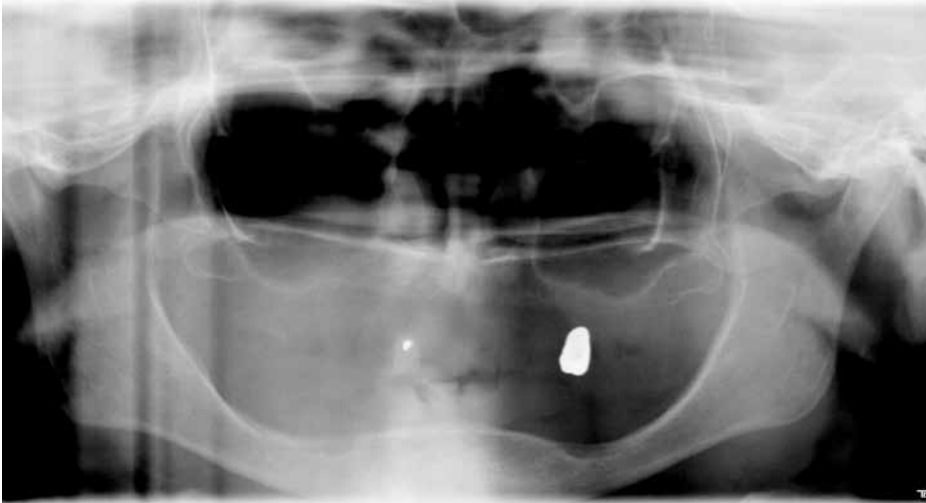


Fig. 1: The severely resorbed mandible before implant therapy.



Fig. 2 : 8 years postoperative radiograph showing the implants and the prosthetic restoration. No vertical (crater-like) bone loss is observed.

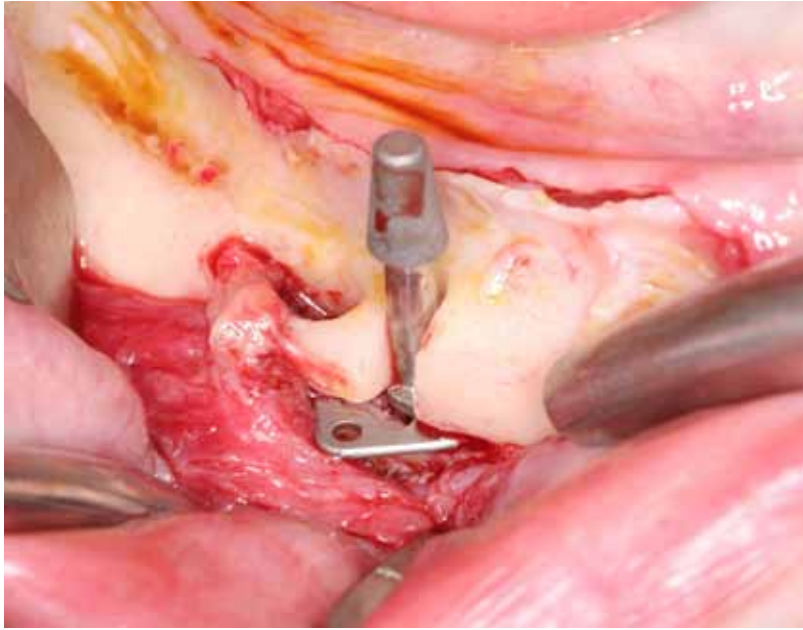


Fig 3: Insertion of a basal implant with the base plate positioned below the mental nerve.

Evidence Report

A comparison of bone grafts with and without platelet-rich plasma in preparation for dental implant placement

Evidence Report Purpose

Long-term success criteria for assessment of dental implants includes radiographic measurement of marginal bone loss. Serial conventional intraoral radiographs have been used to assess changes in bone height. However, a limitation of this method is the inability to detect small changes in bone quality and quantity. Recently, there has been considerable interest in using digital radiography as a means of evaluating and quantifying changes in alveolar bone mass. Digital radiography has been reported to improve the diagnostic capacity of radiographic techniques and to detect small changes in bone density.

Objective

To critically summarize the recently published literature evaluating digital compared to conventional radiographic techniques to assess marginal bone levels around endosseous dental implants.

Summary

One study reported significantly greater levels of marginal bone height when assessed with conventional digitized radiographs compared to digital subtraction images. Further, for digital subtraction images, marginal bone height was greatest for linear digital subtraction images, followed by the enhanced logarithmic scale, and then logarithmic digital subtraction

images. However, the outcome of bone height was not appropriate to assess superiority of radiographic technique. Another study reported changes in marginal bone density to be the greatest for logarithmic digital subtraction images, followed by linear digital subtraction images, and then conventional digitized radiographs. The greater changes in marginal bone density were indicative of the more sensitive radiographic techniques. One other study found agreement in marginal bone height levels between conventional radiography and detailed narrow beam radiography for 61.2% of the observations. Studies were of moderate quality so conclusions based on reported differences should be considered with caution. Additional methodologically rigorous comparative studies with comparable characteristics between groups and longer follow-up are needed to better compare digital vs. conventional radiography methods for assessment of marginal bone levels around endosseous dental implants. One study suggested that logarithmic digital subtraction images were most superior, followed by linear digital subtraction images, to assess changes in marginal bone density around dental implants. However, the reported outcomes varied in all studies and consequently were not comparable between studies

Sampling

A MEDLINE search was performed to identify recent studies published between January 2003 and October 2009 examining bone height and bone density around dental implants in studies comparing digital vs. conventional radiographic techniques. Three articles met our criteria, evaluating the treatment comparison of inter-

Table 1. Medline Search Summary

Terms	Hits	Reviewed
Search dental implants OR dental implantation, endosseous [MeSH]	19,516	
Search (dental implants OR dental implantation, endosseous [MeSH]) AND alveolar ridge augmentation AND comparative study, Limits ENGLISH, Human, Literature containing Abstracts	400	3
Bibliographies from existing literature	0	0
Total Reviewed		3

Common Outcome Measures

Marginal bone height

Marginal bone density

Interventions

Radiographic techniques used to evaluate marginal bone levels around dental implants were compared and described as follows

Bittar-Cortez (Clin Oral Impl Res 2006)

- In a follow-up study, 34 patients underwent evaluation of marginal bone density 1 week after dental implant placement and then 4 months later using: a) standardized conventional intraoral radiographs*, and b) digital subtraction images using the EMAGO® software (Oral Diagnostic systems, Amsterdam, the Netherlands).

est, and are included in this report, Table 1. .

Bittar-Cortez (Dentomaxillofacial Radiology 2006)

- As part of a routine follow-up examination in 22 patients, radiographs were obtained 1 week after implant placement and then 4 months later for assessment of marginal bone height using: a) conventional intraoral periapical radiographs*, and b) digital subtraction images manipulated by the EMAGO® software (Oral Diagnostic systems, Amsterdam, the Netherlands). surgery.

Lofthag-Hansen (2003)

- In 40 patients who underwent dental implantation in the lower jaw, 40 implants were randomly selected for examination during a follow-up evaluation. Six oral radiologists and 1 general dentist assessed marginal bone loss using: a) conventional periapical radiography with the paralleling technique, and b) extraoral detailed narrow beam (DNB) radiography with the Scanora® multimodal radiography system (Soredex; Orion Corp., Helinski, Finland).

* Conventional radiographs were scanned into a personal computer and were digitized for comparison with the digital subtraction images.

Table 2. Comparative studies evaluating bone grafts placed with vs. without PRP in preparation for intraoral dental implant placement.

Author (year)	Study Design	Population	Diagnostic Characteristics	Implant Placement			Follow-up (%)	LoE*
Bittar-Cortez, Dentomax Radiology (2006)	Prospective cohort	N=22; Ni=30 female: NR age: NR	Dental implants placed in the upper or lower jaw	Digital extraoral radiography	Digital intraoral radiography	Conventional radiography	4 months: NR†	Moderate
					N=22; Ni=30	N=22; Ni=30		
Bittar-Cortez, Clin Oral Impl Res (2006)	Prospective cohort	N=34; Ni=53 female: NR age: NR	Dental implants placed in the upper or lower jaw		N=34; Ni=53	N=34; Ni=53	4 months: NR†	Moderate
Lofthag-Hansen (2003)	Prospective cohort	N=40; Ni=40 female: 40% mean age: 68 yrs	Branemark dental implants placed in the lower jaw	N=40; Ni=40		N=40; Ni=40	Mean 4.2 years: 83%	Moderate

N=number of subjects; Ni=number of implants

*Level of Evidence (LoE) is based on study design and methods (Very high, High, Moderate, and Poor)

†NR (not reported) = for follow-up rate either not reported or precise follow-up rate could not be determined since the initial number of eligible patients or number lost to follow-up were not provided.

Table 3. Evaluation of articles comparing digital vs. conventional radiographic techniques to assess marginal bone levels around endosseous dental implants.

Study design and methods	Bittar-Cortez Clin Oral Impl Res (2006)	Bittar-Cortez Dentomax Radiology (2006)	Lofthag-Hansen (2003)
1. What type of study design?	Pro-spective cohort	Prospective cohort	Prospective cohort
2. Statement of concealed allocation?*	N/A	N/A	N/A
3. Intention to treat?*	N/A	N/A	N/A
4. Independent or blind assessment?	NO	NO	NO
5. Complete follow-up of >85%?	YES	YES	YES
6. Adequate sample size?	NO	NO	NO
7. Controlling for possible confounding?	NO	NO	NO
LEVEL OF EVIDENCE	Moderate	Moderate	Moderate

* Applies to randomized controlled trials only

Results

Marginal bone height (Figure 1)

- Marginal bone height at 4 months after implant placement was significantly greater for conventional digitized radiographs ($10.3 \pm 1.9\text{mm}$) compared to DSIs ($p < .05$). Further, when comparing DSIs, marginal bone height was greatest for linear DSIs ($9.7 \pm 2.0\text{mm}$), followed by enhanced logarithmic DSI ($9.5 \pm 2.1\text{mm}$), and then logarithmic DSIs ($9.4 \pm 2.1\text{mm}$). Amongst DSIs, the marginal bone height was significantly different between linear DSI and logarithmic DSI ($p < .05$). [Bittar-Cortez, Dentomaxillofacial Radiology]
- There was agreement in marginal bone height levels between periapical radiography and detailed narrow beam (DNB) radiography for 61.2% (303/495) observations. Periapical radiography showed more marginal bone loss than detailed narrow beam (DNB) radiography for 16.6% (82/495) of the observations, while DNB showed more marginal bone loss compared to periapical radiography for 22.2% (110/495) of the observations. [Lofthag-Hansen]

Marginal bone density (Figure 2)

- Changes in marginal bone density between 1 week and 4 months after implant placement were greatest for logarithmic digital subtraction images (DSI; maxilla: 19.4 gray, range 14.1-34; mandible: 27.9 gray, range 15.6-40.2), followed by linear DSIs (maxilla: 17.6 gray, range 13.2-29.5; mandible: 22.1 gray, range 16.8-33.7), and then conventional digitized radiographs (maxilla: 15.1 gray, range

11.5-25.1; mandible: 17.9 gray, range 14.2-29.1). However, the differences between the groups were not statistically significant. [Bittar-Cortez, Clin Oral Impl Res]

Methodological considerations

- Three studies reviewed were prospective cohorts with a rating of moderate (moderate quality cohort studies) level of evidence. No randomized controlled trials or high quality cohort studies were identified in the literature.
- All three studies had sample sizes that were probably not adequate to show a difference between the study groups for some of the outcomes measured.
- The reported outcomes varied in all studies and consequently present a challenge regarding interpretation and comparison of study outcomes. Further, the outcomes reported for one study (Bittar-Cortez, Dentomaxillofacial Radiology) were not appropriate to determine superiority of the assessed radiographic techniques.
- Since multiple implants in the same subject are not statistically independent, either one implant should be chosen per patient or statistical analysis should account for multiple implants per patient. This did not occur in the two studies for which multiple implants were evaluated in the same patient

Only one study reported a follow-up rate. A follow-up rate of $\geq 85\%$ is necessary to ensure valid study results.

References

Studies

Study 1

Bittar-Cortez JA, Passeri LA, Boscolo FN, Haiter-Neto F (2006)

Comparison of hard tissue density changes around implants assessed in digitized conventional radiographs and subtraction images

Clin Oral Impl Res 17:560-64.

Study 2

Bittar-Cortez JA, Passeri LA, Boscolo FN, Haiter-Neto F (2006)

Comparison of peri-implant bone level assessment in digitized conventional radiographs and digital subtraction images

Dentomaxillofacial Radiology 35:258-62.

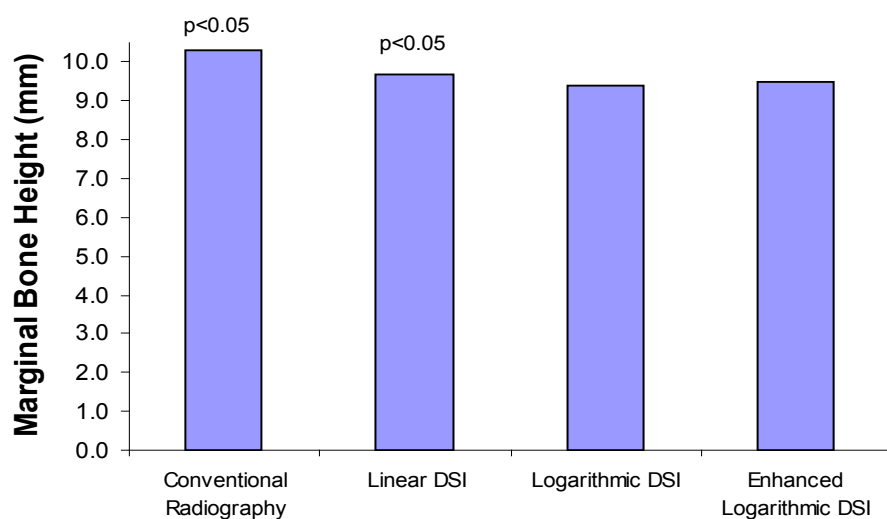
Study 3

Lofthag-Hansen S, Lindh C, Petersson A (2003)

Radiographic assessment of the marginal bone level after implant treatment: a comparison of periapical and Scanora detailed narrow beam radiography

Dentomaxillofacial Radiology 32:97-103.

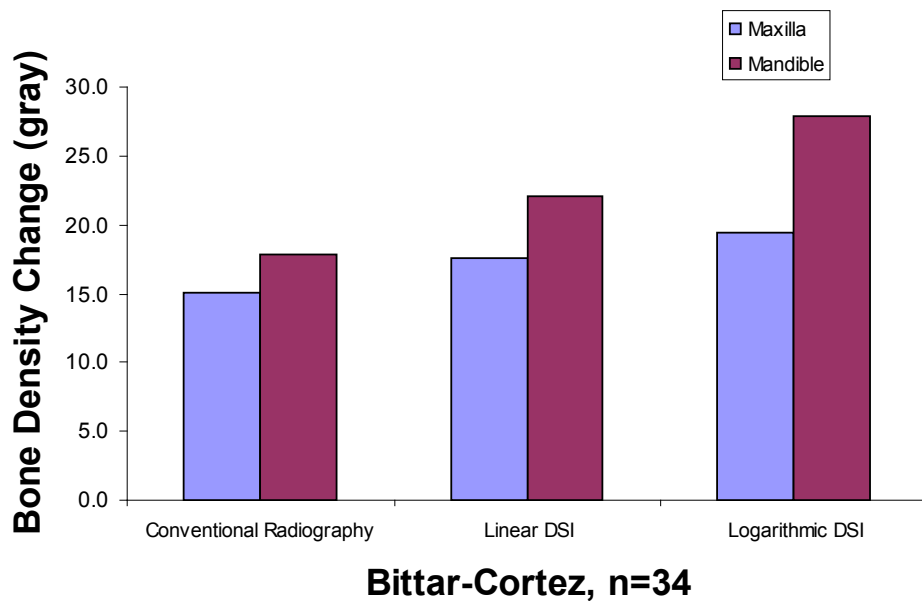
Figure 1. Mean marginal bone height levels using conventional compared to digital radiography methods at 4 months after endosseous dental implant placement.



Bittar-Cortez, n=22

Statistical significance noted on graphs if provided by author

Figure 2. Changes in marginal bone density using conventional compared to digital radiography methods at 4 months after endosseous dental implant placement.



Statistical significance noted on graphs if provided by author

Research in Context

Developing the study plan

Dr. med.dent. S.K.A. Ihde
MNE-85386 Vrba/Tudorovici, Montenegro
Email: ihde@ihde.com

Background: As practicing implantologists, it is important to consider contributing to the current body of implant literature. There are many questions that have either been answered inappropriately or have yet to be answered. Research should not be reserved for universities only. Practicing implantologists have the best ideas and access to patients. Even the best ideas can fail without a good plan. In the last edition of Implant Directions we discussed the importance of the SPECIFIC AIMS. These are the objectives based on the clinical questions that you seek to answer in your study. The specific aims drive the overall study plan.

This Research in Context series is aimed at providing clear and simple guidance on how to go about doing your own study and publishing your results. Though research takes time, it doesn't need to be a second job. With some careful planning, you should be able to publish research out of your own office practice.

The third step in publishing your research is:
DEVELOPING THE STUDY PLAN!

Once the study question and specific aims have been established, you can begin developing your study plan. The study plan is best developed in three stages with each subsequent stage increasing in complexity as outlined below. In this edition of Implant Directions, we will focus on the study outline (Stage 1).

Stage 1	Stage 2	Stage 3
Study Outline	Study Protocol	Study Operations Manual

Stage 1: Study outline or letter of intent (LOI)

The purpose is to provide an outline of the basic elements of the proposed study, and should be one to two pages in length. The basic components for this outline are listed below:

Checklist for LOI.

1	Research Question	√
2	Background and Significance	√
3	Expected outcomes	√
4	Time frame	√
5	Methods and brief research plan -Study design -Subject inclusion and exclusion criteria -Demographic, predictor, and outcome variables -Statistical issues (hypotheses, sample size, basic analysis plan)	√
6	Participants	√
7	Resources and budget	√
8	Research site	√

The study plan and one- to two-page outline should be written at an early stage using the checklist as a guide. Putting ideas on paper is an important step in facilitating discussions and advice from colleagues or mentors. For many, this is a challenge because it is often easier to talk about an idea than to write it down; however, outlining the study will lead to a faster start and improved study. The following table provides an example of a study outline.

Example outline of a study (LOI).

Element	Example
Research Question (s)	<p>Is BOI more effective than conventional screw implants with respect to time to loading, survival, and oral health related quality of life?</p> <p>Is BOI safer than conventional screw implants with respect to complications?</p> <p>Is BOI more cost effective than conventional screw implant systems?</p>
Background and Significance	<ul style="list-style-type: none"> • Immediate loading of dental implants is becoming a critical aspect of treatment decision making. • Patients oral health related quality of life is based on such things as how soon they can chew, potential for pain and other complications, and overall cost of the their treatment. • BOI implants provide a safe and effective alternative to conventional screw implants systems that meet the needs of most patients. • The literature is lacking a study comparing these two treatment methods to establish safety and efficacy.
Expected outcomes	<ul style="list-style-type: none"> • Unknown. • Few comparison studies with sound study methods exist making this comparison. • We hypothesize that patients who receive BOI will load their implants earlier, demonstrate greater levels of quality of life, and spend less on their implant treatment and therapy.
Time frame	<ul style="list-style-type: none"> • 3-month ramp up. • 30-month recruitment, follow-up, and data collection. • 3-month close-out.

<p>Methods and brief research plan</p> <ul style="list-style-type: none"> - Study design - Subject inclusion and exclusion criteria - Demographic, predictor, and outcome variables - Statistical issues (hypotheses, sample size, basic analysis plan) 	<p><u>Design:</u> Multi-site prospective cohort study.</p> <p><u>Subject criteria:</u> Male and female adults aged 18 years and older with all types of bone quantity and quality.</p> <p><u>Predictors:</u> Bone quality, number of implants needed, smoking, other comorbid conditions such as diabetes, length of disability</p> <p><u>Outcomes:</u> Time to loading, survival, complications, oral health related quality of life, and cost</p> <p><u>Statistical issues:</u> Patients will load their implants 3 weeks sooner receiving BOI than conventional screw implants. Rates will be compared using a chi-square analysis and regression methods. Continuous outcomes will be compared using analysis of variance. Estimated sample size (with 80% power) to demonstrate a 3 week difference in time to loading between two groups is 80 (40 per group).</p>
<p>Participants</p>	<p>List co-investigators, collaborators, and sites involved in the study.</p>
<p>Resources and budget</p>	<p>An estimate of costs for study personnel, supplies, subject payments, and consulting costs if applicable.</p>
<p>Research site</p>	<p>Explanation of institution(s) where patients will be recruited and primary data that will be collected and managed.</p>

In the next edition of Implant Direction we will discuss how to use your evolve your study outline into a full study protocol and study plan to ensure success of your study!

Full Length Article

Functionally related changes to the vertical peri-implant bone after BOI implant insertion in the mandible

Author:

Dr. med.dent. S.K.A. Ihde

MNE-85386 Vrba/Tudorovici, Montenegro

Email: ihde@ihde.com

Patients who receive enossally supported bone/implant/restoration systems will experience changes to essential parameters of masticatory function as normal function is restored. Within the jaws, the levels of tension between the affected bone areas will change as well. BOI implants facilitate the enossal transmission of masticatory forces via the baseplates while avoiding the transmission of forces via the vertical aspect of the implant. Consequently, the crestal bone level is able to develop freely to match the masticatory forces, the distribution of functional loads and the tension within the affected jaw. Unlike crestal implants, BOI implants are not associated with any progressive bone loss over time and do not trigger any crater-shaped areas of collapsed bone. In the region of the threaded pin, the bone level can float freely to accommodate the prevailing functional requirements BOI implants were designed to allow these morphological changes. But given that the prosthetic superstructure is the only fixed point of reference within the system, the elevation of the basal disk relative to the mandibular border and masticatory plane requires regular timely subtractive or additive modifications to the fixed restorations in order to protect the

system from prosthetic-related overload and damage to the load-transmitting regions. The extent of vertical bone modelling in the mandible will depend on the preoperative vertical bone supply, the differences between the respective bone mass on either side of the jaw and from the presence or absence of a preferred side for mastication.

Key words

Basal osseointegration (BOI), reversal of mandibular atrophy, functional implantology

Introduction

From a structural point of view, the mandible is essentially a long tubular bone. The chin is an area of maximum flexion in which trajectories from both sides overlap to form a structure of high density. This area is a "safe haven" for implant anchorage, and hence a preferred site for implant placement. The shape of the mandible is derived from a number of factors. Crucial determinants include muscle function (i.e. chewing and mimic muscles) as well as the number, size and position of teeth. When teeth are extracted, the related structure of the alveolar ridge will undergo disuse atrophy, i.e. the vertical bone volume will be reduced in a clinically manifest manner. The delivery of restorations insufficiently supported by residual teeth or enossal anchorage may accelerate bone resorption, influencing the vertical dimension of occlusion and, hence, the vertical dimension of the face at large. Similar to the way in which we differentiate between load-bearing and non-load-bearing structures in buildings, we can

make an analogous distinction in the jaws. The load-bearing structures of the jaws are those areas that possess a trajectory orientation as a result of masticatory function and are designed to prevent mechanical failure of the jaw as a whole. These structures are resistant to resorption. Bone areas whose principal task is to accommodate the dental roots, by contrast, have no load-bearing function. The distinction is not always clear, however. For example, the basal root segments of lower second molars are frequently located in load-bearing jaw segments – which is good for bone preservation (and thus for the preservation of the teeth), if appropriate functional patterns are present.

There have been very sporadic indications in the literature that BOI-based implant/restoration systems not only arrest the bone loss brought about by periodontal disease but can also generate new bone tissue. Follow-up examinations of implant/restoration systems in heavily resorbed areas of the distal mandible have sometimes revealed considerable new bone formation below and above the load-transmitting disks of BOI implants (Figures 9 and 10). To investigate whether these are coincidental findings or a common reaction on the part of the jawbone, orthopantomographs (OPGs) were used to check vertical bone heights pre- and postoperatively as well as after up to 28 months in situ under full masticatory load. Early vertical bone gain following implant insertion and immediate loading was also found in animal experiments, where there was modelling-type vertical bone growth that originated in the periosteal tissue. The scope of the bone modelling exceeded the original level that was

readily discernible on histological sections.

Long-term success can only be achieved if the treatment concept duly considers the functional requirements.

Materials and methods

The target parameter was the vertical bone height around BOI implants inserted at the mandibular first and second molar sites (36/37 and 46/47) placed at our clinic between 1 December 1999 and 1 June 2000 and restored with fixed prosthetic dentures. The place of insertion for the BOI implants varied slightly along the sagittal plane, depending on the prevailing political situation, but was always within the terminal segment of the radiologically documented mandibular linea obliqua.

It was not possible to include implants in the follow-up whose baseline OPG had not been taken digitally by our OPG device (14 affected implants). These OPGs had been taken at other centres and provided to us by the patients themselves. We also excluded those implants that had been inserted in single-tooth gaps for placement of the first molar only (4 affected implants). Three implants had been removed and replaced during the observation period. To the extent that the load-transmitting disks of the replaced (original) implant had remained within the bone at the follow-ups, the measurements were taken at the measuring points of these disks. In the region around one implant, bone growth was so pronounced that the newly formed crestal bone had to be removed by a reductive osteotomy. This implant was not included in the analysis. Overall, 120 BOI implants were available for the follow-up.

These 120 implants (57 of the right side, 63 on the left side) had been inserted in 81 patients (29 male, 52 female). Their average age at the time of placement was 61.5 (29–80) years.

The selected interval for this study corresponded to the first 18 months of use of the digital x-ray system that had been instrumental in facilitating a comparative evaluation of the images with sufficient precision and speed in the first place. All implants have been in function for at least 325 days (0.89 years) and at most 1,050 days (2.8 years) – mean: 674 days – between the first and last radiological examination included in the analysis.

The radiographs were taken with a Orthoslice 1000 OPG system (Trophy, Kehl, Germany); the images were processed and measurements were made using the corresponding 4.1K imaging software. This work was performed on a flat-screen monitor (Dell Inspiron 8200), as it had turned out that measurements on a conventional cathode-ray tube showed much greater variation due to variable viewing angles. All measurements were performed by the same operator by clicking on the relevant measurement points

Measuring points (Figures 7 and 8):

- The upper measuring point was the topmost discernible point of the alveolar ridge at the centre of the threaded pin.
- The middle measuring point was the transition from the bar to the thread holder, i.e., at the level of the baseplate.

- The lower measuring point was located on a line perpendicular to the lower edge of the mandibular cortical bone.

- If the BOI implants had not been placed at the thinnest (vertical) point of the mandible, the vertical dimensions of these areas were also measured.

- If natural teeth were present on the contralateral side of the implant site, the upper measuring point was the intersection of the mesial root of the second molar (or the distal root of the first molar, as appropriate) with the alveolar ridge. This measurement was taken to determine the contralateral bone height for comparison. The vertical distance to the lower edge of the mandible was determined for these measuring points, too.

- Only single-disk BOI implants were placed.

The average preoperative bone supply at the implant site was 19.3 mm (9.9–30.2 mm) for the right mandible and 18 mm (8.9–30 mm) for the left mandible.

All radiographs were taken as part of routine follow-up examinations. No additional patient x-rays were taken for the purposes of this study.

To evaluate their reproducibility of the x-ray images, vertical measuring pins of a known standard length were placed on the vestibular side of the mandibular arches of 10 patients during the x-ray process. These served to determine the discrepancies between the real length and the measured length in the x-ray image as well as the reproducibility of the measurements. The t test showed the error to be non-significant for 90% of the measurements.

Results.

Vertical bone growth in the right mandible was a mean 0.7 mm (2.8–0.1 mm) (Figure 10, areas A and C) above the implant baseplate and a mean 0.9 mm (4.3–0.1 mm) (Figure 10, area B) below the implant baseplate, i.e. toward the mandibular border.

Vertical bone growth in the left mandible was a mean 0.7 mm (5.4–0.1 mm) (Figure 10, areas A and C) above the implant baseplate and a mean 0.7 mm (4.5–0.1 mm) (Figure 10, area B) below the implant baseplate.

This means that vertical bone growth at the implant site was present in all cases during the postoperative observation period. The mean growth was 1.9 mm (6.2–0.2 mm) (Figure 1).

Differences became evident when the changes in the vertical bone levels at the implant sites are related to the original bone height. The greatest amount of bone apposition was observed where the initial overall bone supply at the implant site was minimal while the vertical bone supply in the distal mandible on the contralateral side was massive (Figure 2)

The findings for two patients were clearly at variance with this overall trend in that the amount of bone apposition by the end of the observation period was even greater than the preoperative vertical bone height on the contralateral side.

Prior to implant insertion, 36 patients had worn removable – in some cases tissue-supported – dentures, while 45 patients had either not worn any restorations or received their implants directly at the time of tooth extraction.

Not surprisingly, the group of patients who had not been wearing restorations exhibited a greater initial total bone height of the implant site. The amount of vertical bone gain was greater for the group that had previously worn dentures.

Taken into account the difference between the implant site and the contralateral side of the mandible, patients with considerably more bone volume on the contralateral side compared to the implant site exhibited greater bone gain in patients whose vertical bone levels tended to be done from both sides of the distal mandible (Figure 6). In other words, the differences in bone volume between the two quadrants are balanced.

The pontic regions of 12 patients exhibited areas of lower vertical bone height than the implant sites themselves. These pontic regions benefited noticeably more in terms of vertical bone gain than the implant sites themselves. The mean vertical bone gain in this region was 6.1 mm (Figure 9).

Discussion

BOI implants are successfully employed to provide enossal anchorage for fixed prosthetic dentures. They have greatly expanded the range of indications for fixed restorations, as they can be inserted even in minimal-height residual bone, on condition that the maximum width of the bone is included.

BOI implants can be used in combination with natural pontics or other implant pontics, but they may also be splinted among themselves by fixed restorations.

The vertical component of a BOI implant (i.e. the threaded pin) is smooth and does not show any surface-enlarging features. The surface of the baseplates, however, is usually roughened by sandblasting to maximize the available surface (hybrid design). The masticatory load is transmitted exclusively through these baseplates. While a continuous vertical bone loss has been described for crestal implants in function, blade implants tend to sink into the mandible unless they receive lingual support by the cortical bone. No pertinent results have yet been published for BOI implants.

Bone loss in the region of the threaded pins of BOI implants may be caused by intraosseous infection descending toward the disk or by morphological changes in the jaw caused by atrophic processes. No cases of intraosseous infection were found during the follow-ups in this study. There were only two cases in which the vertical bone gain was so pronounced that a considerable part of the originally intraoral aspect of the threaded pin and of the crown restoration were increasingly covered by bone and overlying because. These regions thus became inaccessible to oral-hygiene efforts, resulting in recurring infection and ultimately requiring re-implantation. In both these cases, the situation during the initial implantation was such that insufficient vertical bone was present at the preferred implant sites (37/47) above the inferior alveolar nerve; the available bone supply was less than 2 mm. This is why areas

further distally on the ascending ramus had to be used instead. After approximately 15 months, reimplantation was possible in the area of the second molars where a sufficient

amount of vertical bone had grown on the crestal aspect of the mandibular canal.

Two patients in whom mandibular implants were placed symmetrically on the left and right sides revealed patterns of bone growth that were essentially unilateral. The preferred sides here were those where the vertical dimension had been greater before implantation. The apposition pattern in these patients was the exact opposite of the pattern seen in all other patients studied.

The results on clinical examination confirmed the finding obtained by interviews with these patients, namely that those masticatory patterns were purely unilateral in nature, as TMJ function to the contralateral side was almost completely obstructed. Vertical bone gain was observed mainly on the non-working side.

This apposition behaviour had been described previously by Hylander. It had been attributed to the fact that compressive forces develop on the non-working side of the lower mandibular margin during mastication, whereas tensile forces are predominant in the upper portion of the alveolar ridge. The results that Hylander had obtained in monkeys were later confirmed in more detail in studies on humans performed by Korioth and Hannam.

To compensate for greater chewing forces, the jaw may grow not only vertically but also laterally or medially. However, these dimensional changes were outside the scope of this specific follow-up study.

Likewise, bone is capable of functional adaptation by changing its degree of mineralization.

Research in crestal implantology is focused on achieving “osseointegration” and ignores the fact that major solid bodies integrated into bone segments that are subject to natural flexion and whose flexural behaviour is different from that of bone will either not be osseointegrated over their entire surface or the degree of bone mineralization will vary along the interface. The human body lacks true reference points. Enossally anchored (osseointegrated) BOI implants can change their position relative to the radiographically visible boundaries of the jaw as the bone morphology changes due to functional influences. Baseplates inserted in the distal mandible tend to migrate upwards in a cranial direction relative to the lower edge of the mandible. This upward migration is theoretically opposed to the chewing force, whose tendency is to push the disk towards the caudal aspect. It appears that the steady force of trajectorial remodelling is stronger than the periodic influence of mastication. Installing force-transmitting surfaces in the enossal space and immobilizing them relative to each other by prosthetic means will create relative reference points. On the one hand, the mastication surfaces guide the muscles; on the other hand, the muscle function thus modified has an effect on the trajectory architecture of the bone structure and implant bed.

The bone must therefore be given an opportunity to change its morphology to adapt and to re-orient its trajectory. The growth tendencies in the crestal direction must not be thwarted by the presence of pontics. In our experience, increasing contact between the pontics and the mucosa may gradually give rise to pain

(frequently at a subconscious level) and avoidance patterns. This in turn may prevent uniform mastication, jeopardizing the integration of implants, or give rise to excessive loading.

In addition, contact between pontics and mucosa will keep the bony structure of the jaws from developing their most favourable biomechanical shape. This, too, may cause problems in the implant bed. For example, growth may occur in directions other than the biomechanically favourable crestal direction (e.g. in width). In the worst case, the load-transmitting surfaces may lose their cortical contact. Since morphological changes of this type can never be excluded with certainty, BOI implants should, if possible, tend to project from the cortical bone rather than being located more deeply.

In cases of advanced distal ridge resorption with sufficient bone volume in the anterior segments to retain a fixed restoration, one should consider not inserting the restoration right away but to adopt a wait-and-see strategy following distal implant placement until the upward migration of the implant associated with vertical bone formation has been completed. Bone apposition might be greater if no masticatory forces are present to counteract it. Furthermore, this would theoretically reduce the need for subtractive adjustments to the restoration. The counter argument would be that it is after the restoration of the distal occlusal surfaces (at the centre of the masticatory forces) that a “normal” functional relationship is created in the first place. As proper masticatory function is restored, further substantive modelling forces can be expected to affect the bone. Even the insertion of a new complete

denture will change the occlusal situation; similar adaptive processes occur after the delivery of enossally supported restorations.

If the lingual aspect of the residual ridge is very high but narrow preoperatively, the threaded pin can be placed laterally to the ridge. This will create roughly the same spatial relationship as if the ridge were growing in a strictly cranial direction after implant placement (Figure 3). In these situations, our approach is to disrupt the trajectories of the residual ridge from the alveolar crest down to the basal mandibular segment by means of a vertical osteotomy created with a tungsten carbide cutter mesial to the implant site. This increases the functional load on the implant osteotomy, which, in our experience, will accelerate healing, improving the chances of successful implant integration. If this osteotomy is not performed, the risk of integration problems is higher because the bone will lack the functional stimulus to close the lateral osteotomy.

The functionally induced build-up of vertical bone in the mandible has also been discussed in connection with transmandibular implants (TMI). This type of build-up appears to depend on the presence of greater chewing forces acting on the bone from an enossal direction as a result of restored function and on the absence of infection-related bone collapses where the implants penetrate the mucosa. Therefore, the transmucosal vertical implant surfaces must not participate in load transmission; also, they have to be polished to high gloss. Thus, while TMI and BOI implants rely on the same principle for load transmission, the handling of BOI implants is simpler.

For several reasons, patients who have previously worn a denture will benefit more from new bone formation than patients who had not previously worn a restoration in the edentulous area. For one thing, the initial situation is less favourable as the disuse atrophy of the edentulous ridge is compounded by denture-induced resorption. Apparently, however, the atrophy is partially reversible. Moreover, chewing forces will decrease in denture patients as they avoid pressure-related pain. The chewing forces can therefore be expected to increase in these patients after a fixed restoration has been inserted. Once the denture is no longer worn, the compressive forces acting on the mucosa – that otherwise cause chronic ischemia of the crestal mucoperiosteal tissue – are reduced. The blood supply through the central mandibular artery is reduced in advanced mandibular resorption.

Atkinson and co-workers demonstrated different regional bone densities in pigs, dogs and humans. Following tooth extraction, bone density in areas exceeding the alveoli would change. In the initial phase, the density would decrease due to a greater degree of porosity. The tunnelling secondary osteons (BMUs) would induce bone remodelling even in the areas surrounding the alveoli. After initial healing, resorption was observed in the area of the extraction socket. Resorption was delayed if crestal implants had been inserted in the area of the alveolus, and bone density was increased; in fact, bone density would sometimes even rise above the baseline level. Nine months after implantation, however, resorption processes adversely affecting the preservation of alveolar bone were

seen in the implant area. On balance, however, the experiments performed by Atkinson and co-workers yielded no evidence of any changes on the lingual side of the mandible following insertion of crestal implants, which is in keeping with our own previous findings with healed but unloaded dental implants in dogs.

Both tooth extraction and the insertion of metal implants will stimulate bone remodelling. Tooth extraction gives rise to disuse atrophy, which the implantation procedure is designed to avoid.

Following the extraction of mandibular molars, the load-transmitting line of the mandible, which usually runs along the vestibular aspect, may be repositioned to the lingual aspect and take a cranial rather than lateral orientation. In dentate patients, the molars prevent the force-transmitting line from establishing its ideal mesiodistal orientation. In patients with deep mandibular periodontitis, the force-transmitting lines become dislocated despite the presence of teeth, which will ultimately cause those teeth to be lost.

Nonetheless, the principal force-transmitting structure of the mandibular bone will remain unchanged if premolars are extracted or replaced by implants.

There are numerous indications in the literature that load-related stress in the mandible (e.g. by muscle attachments and other functional mechanisms) will influence bone growth. Growth is reduced if muscles are removed. By removing them unilaterally, the direction of growth can be modified. Tooth eruption and the sheer presence of teeth will influence bone

remodelling and, hence, the structure of the bone. It has long been known that bone fractures healing in a curved position will straighten over the years. , Frost postulated that the reaction of the bone does not result primarily from compressive and tensile forces but from the tendency of the force acting on the bone to change its curvature. In his theory of flexural neutralization, Frost he summarized that increasing concavity promotes bone apposition, whereas increasing convexity promotes bone resorption. This may be one reason for the sharp increase in vertical bone volume observed in the distal mandible once a denture is no longer worn.

Standardized measuring templates are attached to the implants themselves or to the superstructures to perform bone level measurements of crestal implants. Images can be taken using standard dental x-ray systems using identical angles and conditions. For the present study, this procedure could not be used because it does not document the mandibular border used as a reference point. An analysis of CT scan images of the mandible might have yielded more accurate results than the procedure actually used. However, this would have required 4 CT scans per patient (preoperatively, postoperatively and at 6–9 and 12–18 months) to obtain the requisite data. In addition, the angled of the CT layers would have to match exactly, a requirement that would have caused considerable problems. Even custom installations to lock the cranium in a fixed position probably does not facilitate recreation of exactly the same position as previously given the extensive morphological changes

(increases the vertical dimension, vertical growth); in addition, caudal apposition is also present, and changes in the angle between the ascending and horizontal mandibular ramus also influenced the result. Exact measurements of the distal mandible cannot be made by CT; in addition, the added cost and radiation exposure levels cannot be justified. The procedure used here is routinely employed for posterior measurements in pre-implantological diagnostics to determine the vertical distance from the inferior alveolar nerve. The measuring balls used in the standard procedure do not indicate any image distortion that may be present. While the measurements in this study were made in millimetres and indicated that such on the pertinent graphs, the fact that x-ray systems have a constant magnification factor must always be taken into account. Discrepancies from the actual height may occur, but this error was identified as systematic while ascertaining the reproducibility of the measurements.

The results of this study underscore the extraordinary importance of a competent follow-up protocol for BOI-based implant/restoration systems. Particularly in the first two years following surgery or prosthetic treatment (initial treatment or re-treatment), the relative migration of the basal disk may give rise to significant premature contacts. For this reason, subtractive occlusal adjustments need to be conducted periodically and in a timely manner. The extent of these subtractive adjustments is greater than dentists less familiar with BOI implants would normally expect based on their experience with natural teeth or crestal implants. It is frequently necessary, for example, to extend

subtractive adjustments down into the metal framework. In patients with occlusal deformities, it is usually necessary to rebuild all masticatory surfaces several times as functional blocks are increasingly resolved over the course of treatment. We use bonded composite for this purpose, since this material can be readily added to ceramic surfaces.

Further investigations are needed to examine the morphological changes of the maxillary bone occurring as a result of increasing functional stimuli. Clinical experience has shown that pontics in the maxilla will routinely call for basal adjustment as well. Lever forces pushing the pontics against the mucosa and ridge may cause implant loss.

If the masticatory surfaces following mandibular implant placement are not adjusted in due course, the mandible will suffer damage if a stable natural dentition is present in the maxilla. By contrast, if BOI-based implant-restoration systems in the maxilla are associated with poor bone quality and quantity, damage will more likely be inflicted in the maxilla in the form of overload osteolysis or implant fractures.

In patients with periodontal disease who originally presented with mobile teeth, BOI implant treatment tends to result in continuously increasing masticatory forces, reconfiguration of the occlusal plane, and extensive bone remodelling. In our experience, these patients therefore require especially close monitoring and even more occlusal adjustments than usual.

Summary and conclusion

The shape of the mandibular ridge may change after BOI implants were inserted, involving vertical bone gain both below and above the load-transmitting baseplate.

The bone generated above the baseplate ends up on the lingual side of the edentulous mandible in accordance with the principal trajectories. The formation of new bone mainly takes place to the lingual of the threaded pin.

The extra bone volume routinely formed below the baseplate is capable of raising the implant and the prosthetic structure relative to the occlusal plane. It is therefore necessary to perform subtractive occlusal adjustment on the occlusal surfaces at regular short intervals for at least two years following implantation. Decompression of the TMJ (e.g. when the denture is no longer used) or a relative elongation of the mandibular ramus may occur as a reaction to the upward migration of the restorative structure.

Since vertical bone apposition is also expected in the areas between the implants, the pontics should always be designed with some clearance. If contact with the crestal mucosa is established as the bridge is worn, the pontics need to be reduced via a caudal approach, as the endpoint of vertical bone growth is not known.

Studies performed on humans and animals have consistently demonstrated that vertical bone formation indeed takes place. . Animal experiments have additionally shown that lingual bone apposition also takes place,

possibly as a result of osteotomy-related plastic remodelling of the mandibular ridge for curvature and bone volume compensation.

Since modifications to chewing function must be expected after any changes to the restoration, these changes will also involve morphological adaptations of the bone. This, in turn, may have consequences for the position and loading of BOI implants. BOI-based implant/restoration systems will stabilize within 1 to 3 years. The type and extent of morphological changes during this period will depend on the difference between preoperative and postoperative function and on whether a removable restoration had been worn previously

Since the regular use of the prosthetic structures (stable, symmetrical chewing function) influences bone formation, it is reasonable to assume that prosthetic modifications performed on implants already healed will have far-reaching consequences in terms of bone morphology. Therefore, patients who have received new restorations on existing implants need to be followed up closely by a competent implantologist for another two years.

Generally speaking, a threaded pin of maximum length should be selected for BOI implants, based on the interocclusal dimensions. This will prevent the restoration-bone or restoration-mucosa interface from being overgrown by bone or mucosa even if vertical bone apposition is extensive.

If the vertical bone supply at the site of the second molar is inadequate, an implant is first placed in the lower segment of the ascending ramus

, followed by inserting a fixed restoration to stimulate functional bone apposition. The distal implant can later be replaced with a BOI implant located further anteriorly. The load-transmitting disk of the first implant is normally left in place with this strategy.

Thanks to BOI implants, fixed restorations can even be inserted in situations of extreme ridge resorption. However, the surgical and prosthetic measures required for this purpose will interfere substantially with the biomechanics of the masticatory apparatus. Due to the extensive morphological changes in the jawbone, adjustments to the prosthetic restoration need to be made on a regular basis.

Implant/restoration systems based on BOI implants presumably differ from those based on crestal implants with regard to the nature and extent of the required follow-up protocol. Good patient compliance is indispensable if a stable outcome is to be achieved.

Figures

The dotted lines on all graphs indicate a 85% confidence interval.

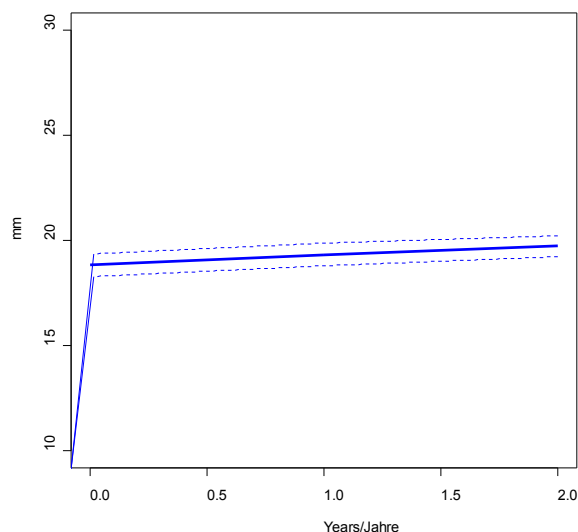


Figure 1

Changes in overall vertical bone height at the implant sites (36/37 and 46/47)

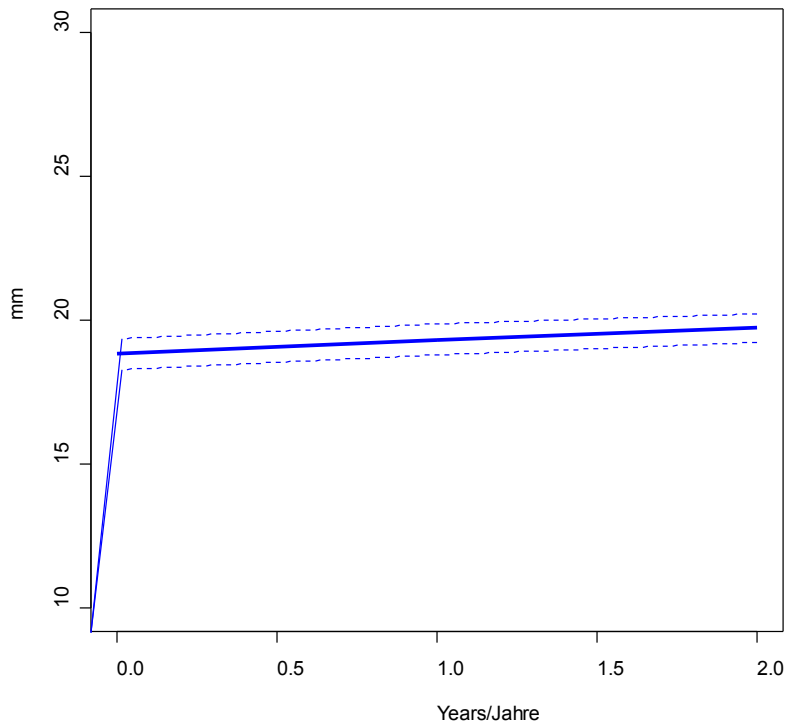


Figure 2

Changes in the overall vertical bone height at the implant sites relative to the initial bone height. For this analysis, each of the original bone height values was assigned to one of four groups: 10–16 mm (blue line; n = 20), 16.1–19 mm (red line; n = 28), 19.1–23 mm (green line; n = 36) and > 23mm (brown line; n = 36)

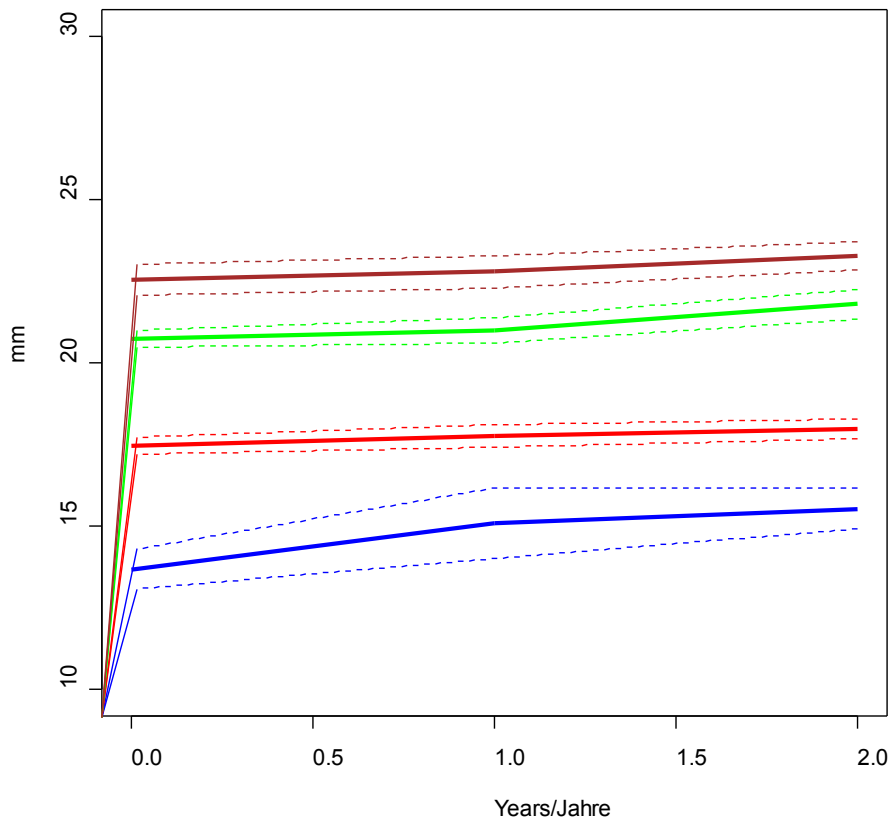


Figure 3

Insertion of a BOI implant at site 47: With BOI implants, the load-transmitting baseplates are inserted in broad basal areas of the mandible; the threaded pin may, if necessary, be positioned to the lateral of any residual thin remnants of the alveolar ridge. These remnants are load-transmitting trajectories that should be interrupted to the mesial of the implant site (light-blue dot/dash line) to improve the chances of healing of the implant osteotomy.



Figure 4

Double BOI implant with a splinted crown restoration: The old cortical bone line is clearly visible. Lingually, there is new formation of lamellar bone. The connective tissue in the shaft area extends below the crown, and there is no epithelial growth in a caudal direction. The soft-tissue structures around BOI implants do not seem to have a layered structure (biological width) like the one described with crestal implants. The implant has been sandblasted for a smooth uniform surface.

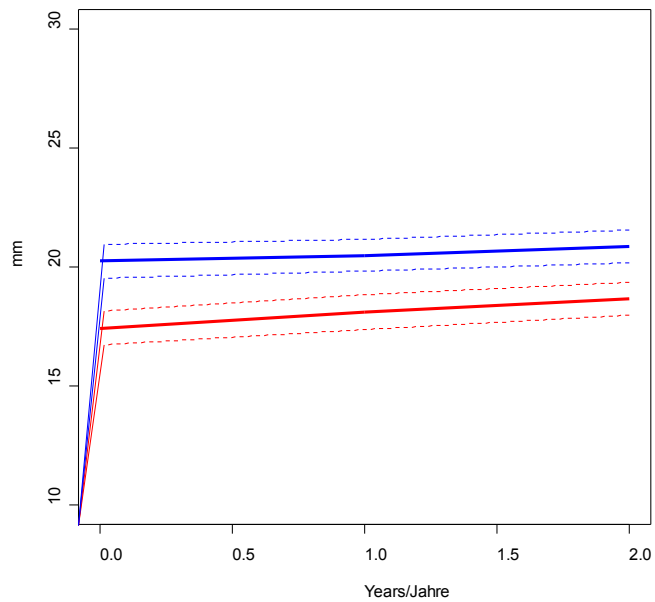


Figure 5

Patients who had been wearing removable dentures prior to BOI implant insertion (red line; n = 36) exhibited more vertical bone gain than patients who had not been wearing removable dentures in the area of the implant site (blue line; n = 45).

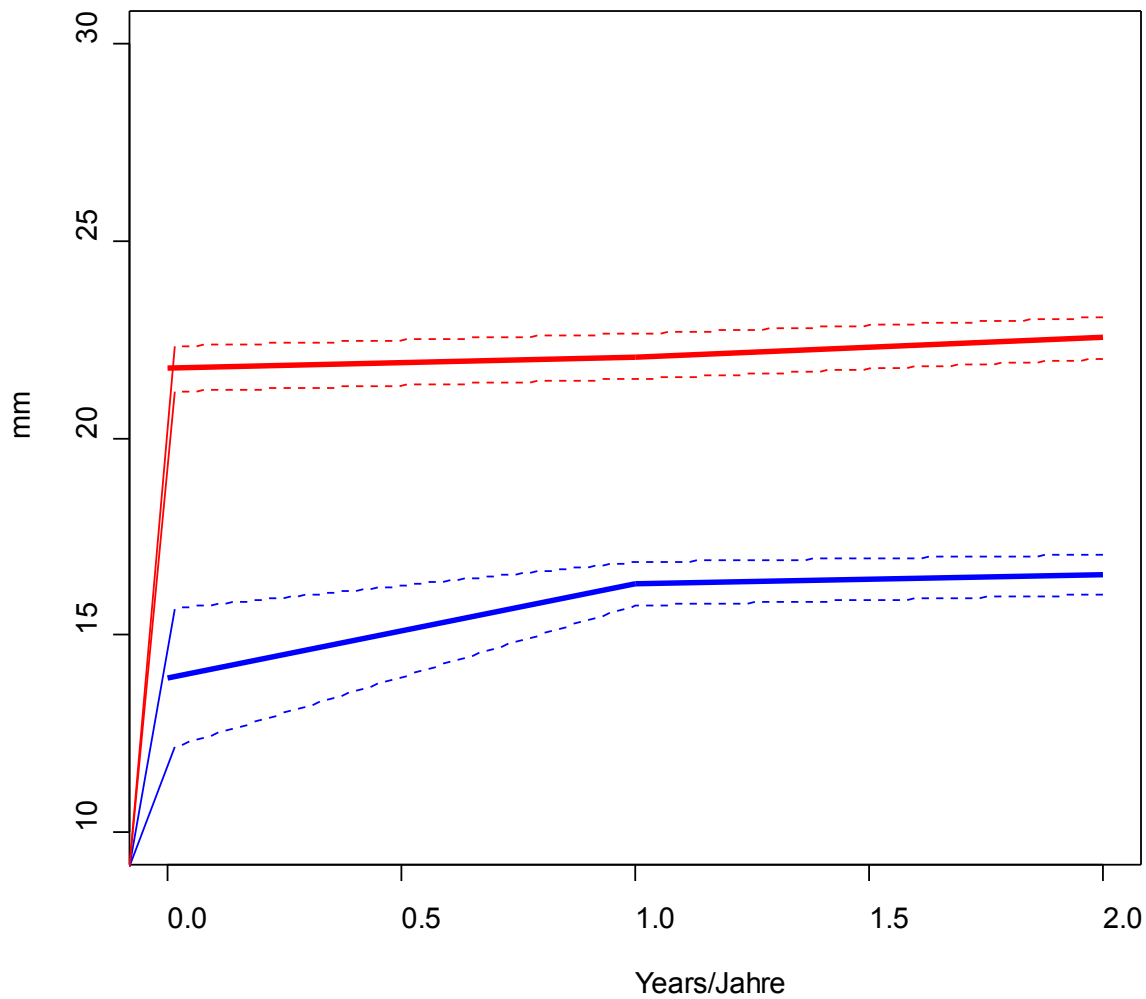


Figure 6

If only minimal vertical bone is present at the implant site that baseline (10–19 mm) and a substantial amount of bone is present on the contralateral side of the distal mandible (> 22 mm), there is more vertical bone gain (blue line; n = 7). If the bone has approximately the same vertical height as the contralateral side of the distal mandible (> 20 mm total bone height on both sides), the amount of vertical bone gain will be less (red line; n = 62).

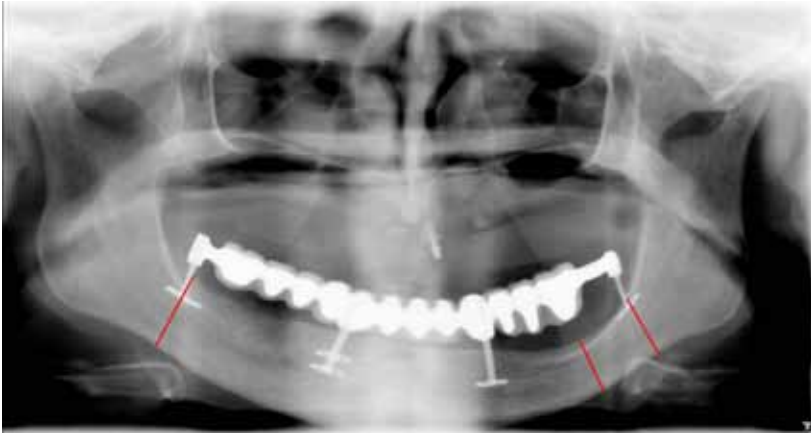


Figure 7

OPG taken after insertion of mandibular BOI implants and delivery of a cemented bridge. This patient had previously been wearing a mandibular complete denture retained only by three ball attachments on the right side. The atrophy of the left posterior mandible had been caused by the denture it had to support.

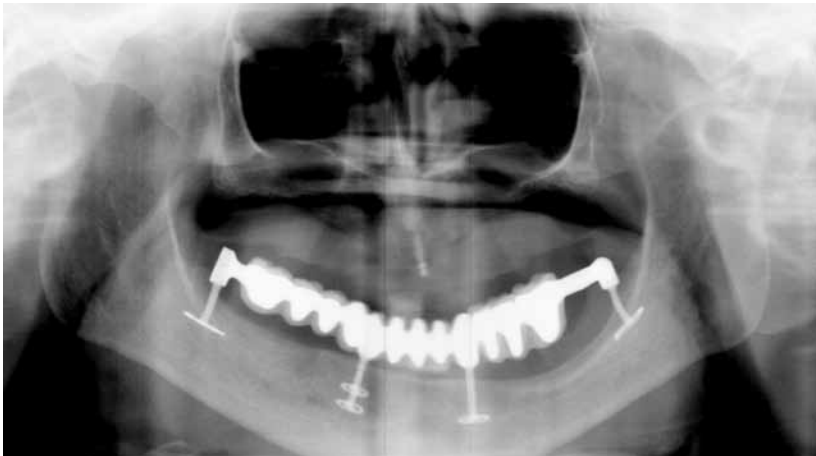


Figure 8

This OPG shows the same patient as in Figure 7, approximately 18 months later. There has been equilibration of bone volume on in the left and right posterior mandible, with a vertical bone gain on the left side of 2 mm. The bone elevations around the extraction sockets still visible in Figure 7 have been levelled. The distance between the ridge and the pontic has been greatly diminished

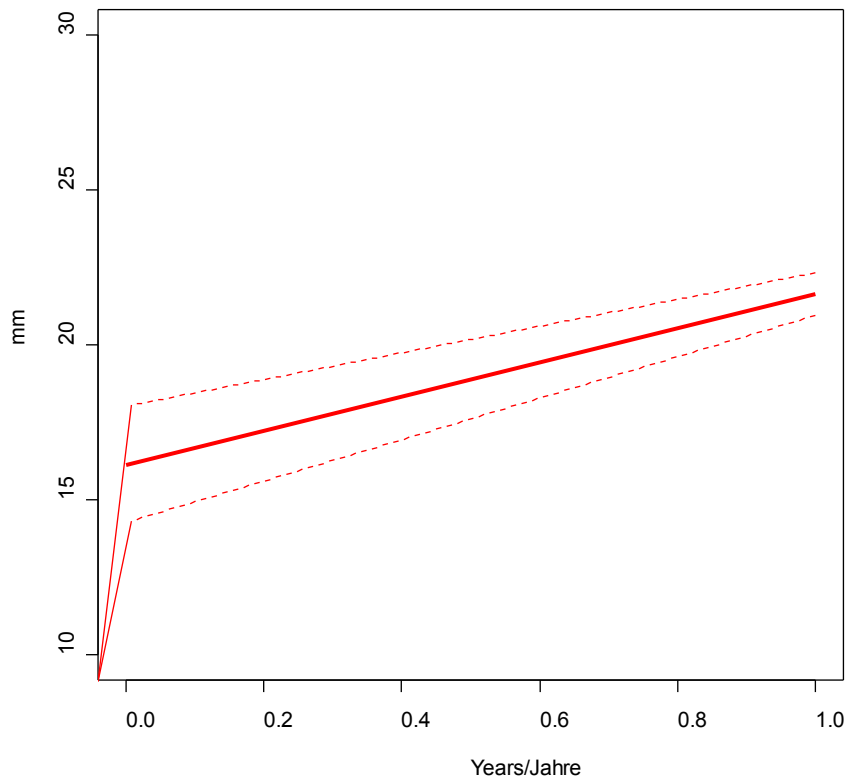


Figure 9
 Changes in total vertical bone height at the lowest points (pontic regions) of the treated mandibles (n = 12).

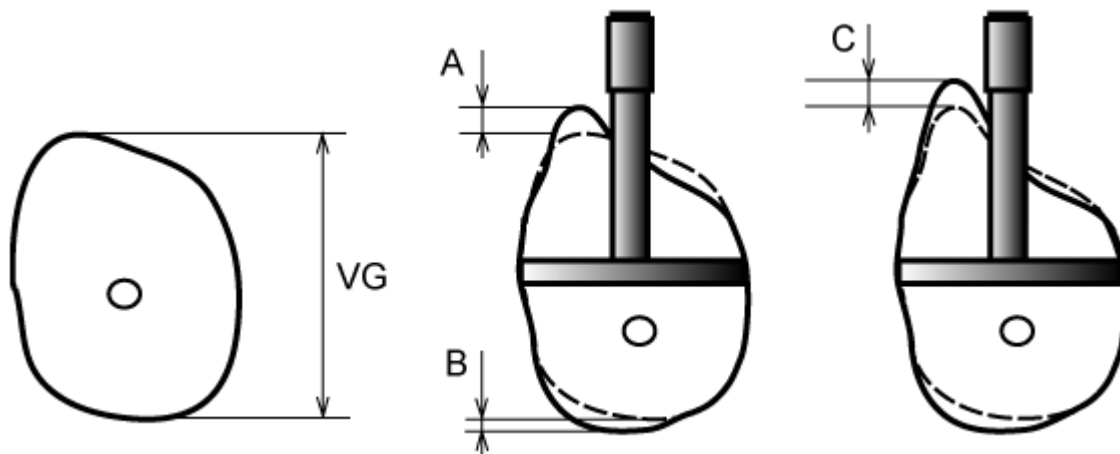


Figure 10

Outline of bone remodelling around a BOI implant in the distal mandible. Left – initial shape of the resorbed mandible. Centre – situation at 12 months after BOI insertion. Right – situation at 24 months. Each dotted line indicates the shape in the adjacent drawing to the left.

- Ashman RB, van Bushkirk WC: The Elastic properties of a Human Mandible. *Adv. Dent Res.* 1(1) 64-67 (October 1987)
- Nähri TO, Etinger RL, Lam EW Radiografic findings, Ridge Resorption and subjective complaints of complete denture patients *The Int. J. of Prosthodontics* Vol 10, No. 2, 1987 (183 – 189)
- Jung F Veränderungen des Prothesenlagers unter der Teilprothese *Deutsch Zahnärztl. Z. Suppl Prothetik und Werkstoffkunde* Jan 1959, Heft 1 8. Jahrg. (105-107)
- Smith BH Changes in occlusal face height with removal partial dentures. *J Prosth. Dent.* Vol 34 No. 3, 1975 (278-285)
- Scortecchi, Misch, Benner: *Implants and Restorative Dentistry*; Verlag Dunitz, London, 2001, S. 428, (ISBN 1-85317-703-2)
- Ihde S., Aleksic Z.: Propagation von Microcracks durch die Sofortbelastung von crestalen und basalen Implantaten; *Deutsche Zahnärztl. Zeitschr.*; in press.
- Ihde S., Konstantinovic V., Cutilo B. Austausch eines BOI unter der festsitzenden Versorgung; *Dent Implantol* 6, 358-361 (2002)
- Schliephake H., Neukam FW Lebenserwartung von Implantaten und Implantatlager *Deutsch Zahnärztl. Z.* 55 (2000) 9 (587-588)
- Hylander WL The functional Significance of the Primate Mandibular Form *J. Morph* 160: 223-240 (1979) sowie: *Patterns of Stress and Strain in the Macaque Mandible* Ann Arbor 1981 Chap 3 und 4
- Korioth TWP, Hannam AG Deformation of the Human Mandible During Simulated Tooth Clenching. *J. Dent Res* 73(1) 56-66 Jan 1994
- Phantarou-Schlieter N.-A.: *Med. Diss Univ. Köln* 2003
- Bosker H., Jordan RD, Powers MO, van Pelt AWJ: Bone induction and bone loss by use of the TMI. *Oral Surg Oral Diagn* 2: 18-28 (1991)
- Mercier P., Vinet A.: Factors involved in residual alveolar ridge atrophy of the mandible; *J Canad Dent Assn* No. 5 1983 339 - 343
- Bradley C.C. A radiological investigation into the age change of the inferior alveolar artery. *Br. J Oral Surg* 13: 82, 1975
- Atkinson P.J., Woodhead C., Powell K.: The Influence of Remodelling on Mandibular Bone Structure; S.263-293
- Atkinson P.J, Powell K., Woodhead C. Cortical Structure of the Pig Mandible After The Insertion Of Metallic Implants Into Alveolar Bone. *Archs Oral Biol* Vol 22, 383-391 (1977)
- Moore WJ An Experimental Studystudy of the functional components of the growth in the rat mandible *Acta Anat* 85: 378-385 (1973)
- Schumaker GH, Dokladal M Über unterschiedliche Sekundärveränderungen am Schädel als Folge von Kaumuskeldissektionen *Acta Anat* 69: 378-392 (1968)
- Zengo AN, Pawluk RJ, Bassett CAL Stress-induced bioelectric potentials in the dento-alveolar complex *Am J. Orthod.* 64: 17-27 (1973)

Jansen M.. On Bone Formation:Its relation to tension and pressure. Longmans, London, 1920.

Basset C.A.L.: Electrical Effects in Bone. Scientific American 213:18-25 (1965)

Frost H.M.: The laws of bone structure. Thomas, Springfield. 1964

Kee-Deog Kim, Ho-Gul Jeong, Seong-Ho Choi, Eui-Hwan Hwang, Cahng-Seo Park; Effect of Mandibular Positioning on the Preimplant Site Measurement of the Mandible in Reformatted CT; Int. J. Periodotics & Restorative Dentistry; Vol 23, No.2, 177 -183 (2003)

Al-Nawas B. Röntgenbilder in der klinischen Forschung – zukünftig ein Widerspruch? Deutsch Zahnärztl. Z. 57(2002) 6; 327 -329.

Besch K.J.: Konsensus zu BOI; Schweiz. Monatsschr. Zahnmed. 9, 1999.



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The text body (headline, abstract, keywords, article, conclusion), tables and figures should be submitted as separate documents.

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