



# 10-YEAR STUDY CHAMPIONS (R)EVOLUTION®

30/03/2022

The aim of this retrospective study was to quantify the survival rate of endosseous implants of the CHAMPIONS (R)Evolution® type under normal practice conditions in a defined time frame (01/09/2011–01/09/2021) compared to the researched study population of the same subject. Hypothetically, comparability is possible.

## Material and methods

The prerequisites and practice conditions are presented, and functional capacity definitions in situ are developed. They are defined based on the results of the 3<sup>rd</sup> ITI Consensus Conference –August 2003. Using the EXCEL software program, they are evaluated concerning patients and implants.

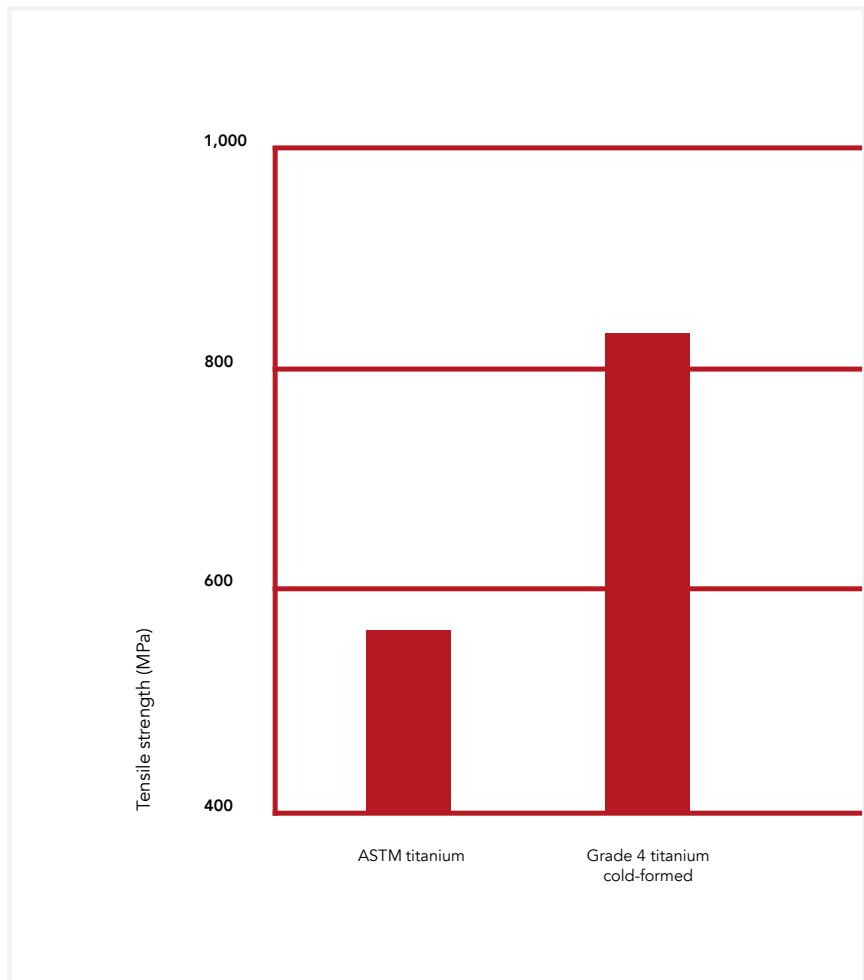
In this study 13,834 Champions (R)Evolution implants were placed in 4,192 patients. Functional capacity in situ, periodontal probe measurements and radiological checks were ideally examined after an observation time of 1 year, 2 years, 5 years, and 10 years.

The national and international literature search was performed via DIMDI and PUBMED according to topic.

Product and service quality, constant innovation, a good price/performance ratio, and sustainability are the four pillars of a successful implant system. Currently, with this 10-year study, the German company Champions-Implants GmbH, with its own full production in Baden-Württemberg, Germany, has now managed to underpin its sustainability and clinical success with this study.

In addition to the titanium one-piece implants available since 2006, the two-piece CHAMPIONS (R)Evolution system has conquered hearts of many implantologists and prosthodontists in Europe since 2011 with over half a million implants sold.

Thus, this innovative, high-quality CHAMPIONS (R)Evolution system, which is "Made in Germany", has been an established implant system at the leading edge.



With the introduction of the system in September 2011, an implant study was started, which was conducted in 4 Implantology dental practices in Europe until September 2021. The aim of this study was to determine the sustainability of this system over 10 years and also to scientifically substantiate its success.

Short introduction to the CHAMPIONS (R)Evolution system, which was designed by Dr. Armin Nedjat based on his previous implant developments ("Classic", "New Art", and Ball-Head) and designs from 2006:



## Material

Like all titanium one-piece CHAMPIONS implant systems, the CHAMPIONS (R)Evolution implant is made from cold-formed ASTM grade 4 titanium, consisting of 99% pure titanium, which is unlike grade 5 titanium (Ti-6Al-4V), containing 6% aluminium and 4% vanadium.

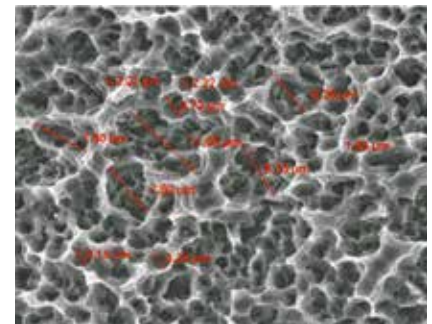
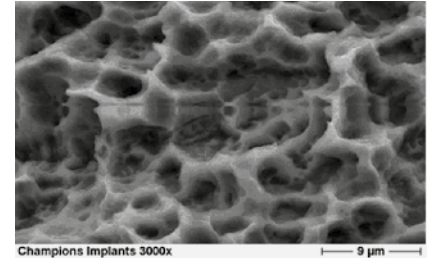
## Implant Design

The main features of the CHAMPIONS (R)Evolution implant are its crestal micro-thread, the internal double cone with integrated Hexadapter (unchanged since 2011 and sufficiently long) and the Shuttle with the mounted Screw ex works, which connects the Shuttle to the implant.

The self-tapping compression thread has a very clean micro-rough surface structure (final testing by CleanImplant). The "implant finishing" is done with irradiation with  $Al_2O_3$  particles afterwards, so that the microrough surface is created in a standardized way. CHAMPIONS is also well positioned with regard to the new MDR.

The Shuttle and Abutment connection with the ideal  $9.5^\circ$  cone prevents bacterial penetration so that the „Zipprich's brine effect" (University of Frankfurt) does not come into play. In 2012, studies conducted by Zipprich showed that the CHAMPIONS (R)Evolution implant had only a maximum micro-gap of  $0.6\mu m$  even with a diameter of 3.5 mm (the smallest bacterium size is about  $2\mu m$ ), while no micro-gap could be verified at all with the larger implant diameters of 4.0, 4.5 and 5.5 mm.

Due to the inner cone and the fact that only one prosthetic platform is available for all implant diameters, which leads to the so-called platform switching effect in the CHAMPIONS (R)Evolution. Platform switching generally means that the implant shoulder is wider than the abutment emergence profile. With an implant with platform switching such as that of the CHAMPIONS (R)Evolution, bone even grows over the implant shoulder and enhances an increase in the stability of the soft tissue cuff, which provides a perfect solution for a periimplantitis prophylaxis.



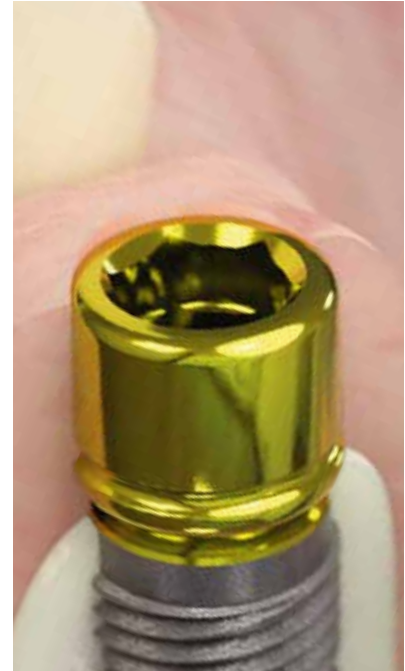
## Shuttle

Currently, the revolutionary so-called Shuttle of the CHAMPIONS (R)Evolution has been a great innovation. It is factory-fitted with a Screw in the implant inner cone, which is also used later for the abutment to fit the denture. The Shuttle fulfills 4 functions. As a rule the Shuttle ensures that the inside of the implant remains sterile until the end of the prosthetic procedure and that the implant inner threads and the implant wall are protected to the maximum during surgery and during various prosthetic processes because they cannot be damaged.

First of all, the Shuttle serves, figuratively speaking, as a "carrier rocket" to bring the "spaceship" (i. e., the implant) safely into the "orbit" (i.e. into the bone) and to keep the inside of the implant sterile without deforming the "thin" outer wall.

Since titanium is a relatively soft metal and the titanium wall at the cone connection is only 0.4mm for a 3.5 mm implant diameter, the Shuttle is a unique tool for avoiding later prosthetic screw loosening since deformation of the thin wall during insertion is excluded. It also serves as the first and only healing abutment since it already has a height and diameter of 3.5mm. However, the Shuttle can still be fitted with a PEEK Gingiva-Clix (preferably after the healing time) for an optimum emergence profile or with two easily prepared PEEK Provi-Clix with different angles directly after implantation with immediate loading. For both Clix, it is not necessary to loosen the Screw or even remove the Shuttle – they are simply set on the Shuttle while the inside of the the implant remains sterile.

There are 6 different Gingiva-Clix allowing for optimum shaping of an emergence profile.



## Shuttle has 4 functions all-in-one

- Insertion Aid
- Surgical Cover Screw
- Healing Abutment
- Impression tool



**Transmucosal impression with a conventional spoon with a Metal Impression Coping (a) and PEEK Coping or with the Impression Coping, Win! PEEK (b)**

Of course, one of the main advantages of the Shuttle is the exact, safe and "closed impression" with the Shuttle and with the help of the Impression Copings, Win! PEEK. Another possibility is also to use metallic Impression Coping (also through the Shuttle, but with removal of the Screw) and the "open impression" in the implant itself (after removal of the Shuttle and the Screw).



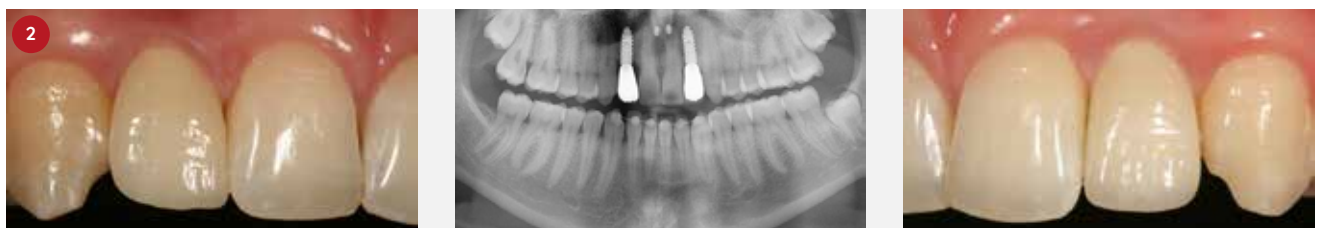
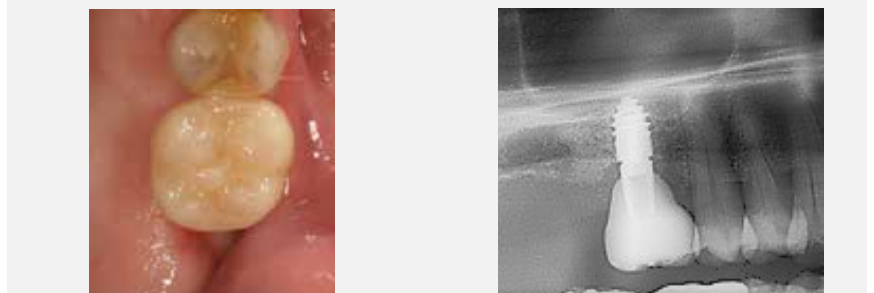
In both cases, a Laboratory Analog for the CHAMPIONS (R)Evolution and a Laboratory Shuttle screwed over it must be mounted to obtain a master cast.

**Indication, Lengths, and Diameters**

The indication range of the CHAMPIONS (R)Evolution implant includes delayed as well as immediate implantation in all jaw regions. Tried and tested, this high-quality „Made in Germany“ is easy-to-use and optimizes the dental workflow of Implantology practices. CHAMPIONS (R)Evolution implants are available in the lengths of 6.5 – 8 – 10 – 12 – 14 – 16 mm and in the diameters of 3.5 – 4.0 – 4.5 and 5.5 mm. Thus, sinus lifts (IDS) and augmentation-accompanying measures are also possible.

**Case report 1:** immediate implantation in site 16 with Champions (R)Evolution implants of Ø 4.5mm and a length of 8mm; use of Smart Grinder; Internal Direct Sinus Lift (IDS).

**Case report 2:** delayed implantation 12 +22 (CHAMPIONS (R)Evolution).



**Prosthodontics**

There is only one prosthetic platform for each CHAMPIONS (R)Evolution diameter, which facilitates (r)evolutionary work in the dental laboratory and in the Champions practices.

Since 2011, all implant abutments have been kept sterile when packaged since 2011 and can therefore be used directly in the mouth. The standard abutments are available in gingiva heights GH: 1 – 2 – 3 – 4 – 5 mm and angles of 0° – 15° – 22.5° – 30°. The standard Abutments are available in gingival heights of GH : 1 – 2 – 3 – 4 – 5 mm and angles of 0° – 15° – 22.5° – 30°.

Abutments Massive are securely fixed in the mouth, prepared with NEM-drills and water cooling as well as shaped like a natural tooth and then cast.

- The dental laboratory is not aware that it is an implant . → cost-saving
- In addition: no impression with transfer posts and no Laboratory Analogs

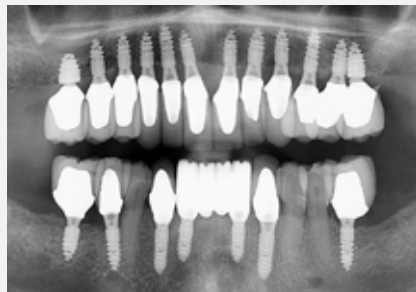
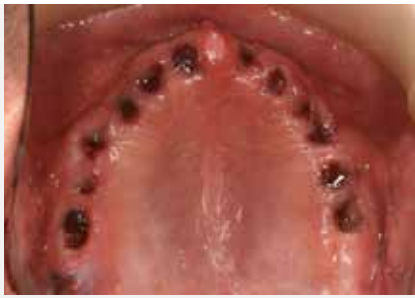
The ICAs = Individual Connecting Abutments are an innovative speciality of Champions-Implants GmbH: for the optimized workflow also in the laboratory, 12 common – already sintered – zircon secondary molds are provided industrially. These can be further individually prepared by the dental laboratory and bonded to the titanium base ICA.

**Case report 3:** ICA-Abutment restoration in site 22

**Case report 4:** immediate implantation in sites 16–26 and 36, 33, 32, 42, 43, 45 and 46 (CHAMPIONS (R)Evolution) with Hybrid Screws Georgi and Smart Grinder



4



Multi-Unit LOC



Titanium base



Calcinable



GH1



GH2



GH3



GH4



GH5



GH1



GH2



GH3



GH4



GH5



GH1



GH2



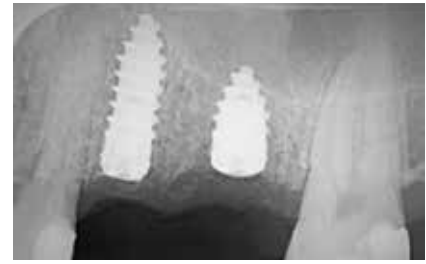
GH3



GH4



GH5



Case report 5: Immediate implantation 25 and delayed immediate implantation in site 26 in surgery and Prosthodontics



**Developer of the CHAMPIONS (R)Evolution system & educational background**

It was in 1995 that Dr. Armin Nedjat developed the MIMI®-method (Minimally Invasive Implantation Method). Then, he perfected it with his Condensers, which he developed in 2006, and the CHAMPIONS (R)Evolution system, which he developed in 2011.

Dr. Nedjat was nominated for the “German Medical Award” in Berlin 2017 and won the SENSES award for “Best Innovation in Medicine” in Dubai in 2013.

With the CHAMPIONS (R)Evolution implant system, the company Champions-Implants GmbH, the biggest German dental implant company, marked a quantum leap forward in Dental Implantology in 2011.

An unconditionally optimized workflow with high-quality, innovative products has facilitated the daily routine in our dental practices since 2011. Due to the useful design concept, high primary stability can be achieved, especially in case of immediate implantations (extraction and implantation in one session). The time and cost savings in dental practices are considerable, in many cases exceeding 100%.

From surgery to Prosthodontics, dental practices have a tried and tested, cost-efficient system at their disposal, which is also fully digitally CAD/CAM capable. The main advantages in today’s digital age are the ability to network with real Implantology experts in almost all languages and guaranteed product availability in the long-term.

The concept of the „Clinical Implantology & Implant Prosthodontics Curriculum“ is based on Prof. Dr. Jean-Pierre Bernard of the University of Geneva. As early as in the 90s he succeeded in teaching Implantology novices skills for incorporating implant treatments into everyday practice through close supervision and clinical training with supervision in each participant’s own practice in Switzerland.

In collaboration with the Future Dental Academy GmbH, the VIP-ZM society (Verbund der innovativ-praktizierenden Zahnmediziner/innen e. V. [German society of dentists with innovative practices who share scientific knowledge]) has successfully introduced this one-year Curriculum in Germany since 2017. More than 150 participants have been trained.



## Prospects of success of implant treatments

Since Greenfield pioneered the cage design of dental implants (Greenfield, 1913), Implantology has been widely disseminated worldwide via the pin implants propagated by Scialom and Pruin (Scialom, 1965; Pruin, 1974) and the blade implants of Linkow (Linkow, 1972). In the 70s, Koch and Kirsch went further with the IMZ system (Kirsch, 1980; Koch, 1976), Schulte with the Tübingen immediate implant (Schulte et al. et al., 1978), and Schröder with his hollow cylinder implants (Schroeder et al., 1976) were a step forward in Implantology. In the mid-1960s, however, Brånemark made important Implantology discoveries. Brånemark and his coworkers worked with a titanium cylinder screw as the basic element (Brånemark et al., 1969). It was Brånemark and his research group who coined the term „osseointegration“, the direct connection between the bone and the implant, which they observed while working with machined, covered, unloaded healing screw implants.

In addition, Schroeder discovered functional ankylosis, a phenomenon of bone healing that he observed with additive-roughened titanium hollow cylinder implants that healed with transmucosal loading. Furthermore, Schulte and his coworkers worked on immediate implantation, which has marked another milestone in the evolution of modern scientific Implantology. The implants that are used today are predominantly rotation-symmetric titanium two-piece screws.

Clinically, Implantology has been growing in importance especially in Germany since DGZMK (Deutsche Gesellschaft für Zahn, Mund – und Kieferheilkunde [German society of Dentistry and Oral Medicine]) officially recognized Implantology as scientific discipline in 1982 (Spiekermann, 1994; Tetsch et al., 1990). Since then, the use of implant-retained dentures in Dentistry has attracted a lot of interest from both the public and the scientific community.

In addition to the stabilization of dentures in an edentulous jaw and the avoidance of dentures especially in

free-end cases, aesthetically demanding areas such as interdental gap restorations or single-tooth dentures have now been routinely performed. In addition to improving chewing function, the therapeutic focus is now often on preserving or regenerating sufficient bone and soft tissue as well as on the preventive preservation of tooth substance by avoiding the reduction of natural teeth.

In recent years, the Implantology field has become very dynamic. In addition to the wider range of indications through various bone replacement graft materials and the modification of implant surfaces, biological growth factors (platelet rich plasma, bone morphogenetic proteins) have been placed, and immediate restoration with dentures or immediate loading has been performed, accompanied by an increased aesthetic demand. Despite a high clinical reliability and predictability of the therapy result, implant failures are still part of everyday clinical life.

Reasons for implant failures can include: Peri-implant infections, connective tissue encapsulation/ failed osseointegration, implant fracture, screw fracture, coating fracture, and iatrogenic reasons (incorrect position, oncology), titanium oxide intolerance, patient psychoses.

Systematic observational clinical studies have long documented the excellent prognosis of endosseous implants through positive long-term results (Adell et al., 1981). The long-term success of osseointegrated implants is the basis of the implants for the established indications (DGZMK, 2000).

The value of implant-retained dentures has been proven due to the high degree of functional restitution even in difficult anatomical conditions (Neukam and Buser, 1996).

The prognosis of osseointegrated implants, however, partly depends on the patient's state of health. Influences of disorders of bone metabolism, diabetes mellitus, radiotherapy, and nicotine abuse on the overall prognosis of implants have been discussed in literature (Blanchaert, 1998).

For instance, it has been believed that osteoporosis (with a bone quality decrease) could be associated with a limited implant prognosis. Although osteoporosis increases with age and after menopause, Dao et al. have shown that implant failure rates are not related to age and gender. A scientific verification of the literature does not provide a compelling theoretical or practical basis to expect osteoporosis to be a risk factor for osseointegrated dental implants (Dao et al., 1993).

Behneke et al. conducted a study consisting of 35 patients with diabetes mellitus who received 129 interforaminal ITI implants in the mandible. With insulin therapy, increased bone resorption was observed during the healing time. A significantly higher resorption with progressive tendency was observed in patients with the underlying disease persisting for more than 20 years compared to patients with shorter disease duration. Within a 5-year observation time, however, there were no differences in the implant-related success or dwell probability between the patients with diabetes (94.9%) and the control group (91.6%) (Behneke et al., 1998). Diabetes mellitus is therefore not considered as a contraindication for implants. The implant prognosis and potential prognostic factors in the irradiated jaw were investigated in a group of 47 patients with 197 implants by an analysis of the implant survival rate (Kaplan-Meier).

Despite the difficult conditions for tumor patients (compliance, hygiene, soft tissue replacement, osteoplasty), the survival rate was 95% after 1 and 2 years respectively and 72% after 6 years. Compared to a historical group, the survival rate of teeth that were healthy before radiotherapy was significantly lower (1 year: 75%; 5 years: 45%) than the prognosis of endosseous implants (1 year: 95%; 5 years: 72%) (Grötz et al., 1999). Despite the significantly reduced prognosis, endosseous implants are helpful in serving as good rehabilitation of the masticatory function also for irradiated patients. Although florid periodontal disease of the residual dentition is considered as a prognostic factor in implant therapy, prospects for this clinically common problem are supported

by little data. A current study by Mengel et al. (Mengel et al., 2001) comparing partially edentulous patients with generalized chronic periodontitis and those with generalized aggressive periodontitis showed a 100% implant success rate in patients with chronic periodontitis after 5 years. By comparison, after this time, the success rates recorded in patients with aggressive periodontitis were only 88.8% (maxilla: 85.7%; mandible: 93.3%). The authors have concluded that implants are a possible solution for these patients. However, according to the authors, as there are no clinical and microbiological differences between the natural teeth and implants, progression of the disease cannot be ruled out. Thus, aggressive periodontitis of the residual dentition should be considered as a risk factor for implant therapy even under study conditions. Nicotine abuse is also known for negative effects on peri-implant hard and soft tissue. Due to wound and healing disorders, smokers experience 3.7 to 7.5 times more failures already in the early phase (Vockner, 2001). Due to the constituents of tobacco smoke such as nicotine, nitrosamines, polycyclic hydrocarbons, benzanthracene, cyanide, heavy metal, and carbon monoxide, the effectiveness of the infection defense is severely impaired. Smoking increases the risk of peri-implantitis by reducing the immune defense. In a study by De Bruyn (452 implants), the differences in implant failures (9% of smokers, 1% of non-smokers) were statistically significant

(De Bruyn and Collaert, 1994). Studies on the influence of nicotine abuse on implant healing and implant prognosis did not show a statistically significant influence of smoking on survival rates of implants in the local bone without additional surgical procedures. Studies on the influence of other systemic diseases on implant prognosis (including collagenosis and arteriosclerosis) are still pending, and in the literature there is only casuistry.

### Evaluation of the implant success

Therapy form evaluation includes a realistic evaluation of the long-term success. Although the implant success evaluation according to the criterion in situ or explanted is clear and can also be determined precisely at the time of the event, different authors consider this sole criterion as insufficient for a differentiated therapy comparison due to the lack of clinical parameters.

As a standard statistical method for the time-dependent investigation of the parameter implant failure, the survival time analysis according to Kaplan and Meier (Kaplan and Meier, 1958) is frequently found in the literature. In this evaluation, the dwell probability of an implant is calculated on the basis of the failure rate over time and the number of implants at risk at that time whereby the implant service time and the number of implants at risk are given a considerable weighting.

However, the Kaplan-Meier dwell

time analysis only considers the fact whether an implant is still in situ after a certain time or not. The condition of the peri-implant soft and hard tissue parameters is not taken into account. However, more and more authors consider the evaluation of these peri-implant hard and soft tissue parameters (Behneke and Behneke, 1996; Moberg et al., 1999), mostly using a Cutler-Ederer analysis.

Corresponding implant success criteria, which include follow-up variables collected during clinical or radiographic follow-ups, have been postulated by various groups of authors (Albrektsson et al., 1986; Buser et al, 1990; Jahn and d'Hoedt, 1992; Naert et al, 1992; Schnitman and Shulman, 1980; Snaauwaert et al, 2000), but a general consensus has not yet been reached.

The average success rates for endosseous implants generally reported in the literature are comparable only to a certain extent due to the use of different implant systems for different indications and because of different evaluation criteria. Data vary between 61% and 98% (Albrektsson et al., 1988; DGZMK, 2000; Dietrich et al., 1993; Richter et al., 1992).

The current analysis of the parameters of a patient group (n = 13,834 implants, 4,192 patients) collected in this study showed implant in situ rate of 95%, which is comparable with the one mentioned in literature or even better, and a Kaplan-Meier dwell prob-

### International criteria for endosseous dental implant success

NIH-Conference, Schnitman 1980	Albrektsson et al, 1986	Buser et al, 1990	Jahn, d'Hoedt, 1992	Naert et al, 1992 – Snaauwaert et al, 2000
Implant in situ	Implant in situ	Implant in situ	Implant in situ	Implant in situ and implant-retained denture
Loosening degree 0–1	Loosening degree 0	Loosening degree 0–1	Loosening degree 0–1	Periotest value < +8
Radiologic, peri-implant translucency is graduated but does not contribute to the definition of success	Absence of peri-implant radio-translucency	Absence of persisting peri-implant radio-translucency	X-rays may not demonstrate any evidence of a bilateral, continuous gap with a width > 0.5 mm in the implant	No peri-implant radio-translucency
Vertical bone loss is not bigger than 1/3 of the vertical implant length	Vertical bone loss < 0.2 mm/annually following the implant's first year of service		The angular bone defect (mean value of the mesial and distal measurement in the X-rays) may not exceed 3/10 of the constructive-endosseous implant section	
Gingivitis accessible to therapy + absence of signs of inflammation	No signs of an infection	No peri-implant infection with putrid secretion	Sulcus depth may not exceed 4 mm for 2 consecutive controls	No implant fracture
No injuries of nerves, teeth, maxillary sinus, or floor of the nasal passage	No injury of nerves	No persisting discomfort such as pain, foreign body sensation, and/or dysesthesia	Subjective evaluation of the implant may not be worse than „3“ (German school grade system)	No implant-induced pain, infections, or paresthesia

ability of 96.5% after 6 years. However, substantial differences are found in the resulting implant success rate depending on the success criterion used:

According to definition of success of Naert et al., the success rate resulted in almost 98%; with criteria according to Albrektsson et al. and Buser et al., the success rate resulted in 97.5%.

The NIH criteria limit annual bone loss and also rate 97.5% as success while Jahn and d'Hoedt see only 95% as success.

An important innovation of the success criterion according to Jahn and d'Hoedt (Jahn and d'Hoedt, 1992) is the inclusion of patient evaluation of the implant according to the German school grade system. For the first time, an only sufficient or even worse evaluation of the implant success by the patient leads to the evaluation of the implant as a failure, which is reflected in the significantly reduced success rate.

In the follow-up examination of the patient group already mentioned above, the patient degree of satisfaction was analyzed on the basis of a 6-grade scale of the German school grade system (6 = dissatisfied, 1 = extremely satisfied); patient degree of satisfaction with implant surgery (MIMI) results was described as good or very satisfactory in 98% of the patients.

95% were very satisfied with the denture and 97% with the total success of the treatment. 98.5% of the patients would be willing to undergo the surgery again if indicated.

99% would recommend the surgery to others. After comparing clinically collected parameters with subjective evaluations of the patients, the Spearman correlation analysis evaluated no statistically significant correlations.

Despite a large number of cases (n = 4,192 patients), the expected influence of clinical parameters on patient satisfaction did not show to be significant.

Only recently has the esthetic success of the restoration been analyzed more and more as a criterion for implant success. Particularly when evaluating immediate loading concepts, it is necessary to make trade-offs in the relevant esthetic zone between early loading and calculable esthetic results with time-tiered forms of therapy. However, the evaluation seems difficult, although this point seems to be integrated in the subjective satisfaction of the patients with the implant restoration, which, however, is overlaid by the different expectations. As early as 1994, the Federal Ministry of Health pointed to patient satisfaction as an essential part of the outcome quality (Anderson, 1998; (BMFG), 1994). In recent years, patient satisfaction has become increasingly important, especially in the context of quality assurance (Leimkühler, 1996).

While as a rule the objective performance is the same for all patients, the subjectively perceived performance can vary individually (Homburg and Rudolph, 1995; Jacob and Bengel, 2000).

Patient satisfaction at the end of treatment plays a significant role in the evaluation of the quality of medical interventions and will continue to gain in importance in the future.

The implant survival probability of which the only success criterion is the implant remaining still in situ at the time of examination regardless of possible complications was analyzed according to the method of Kaplan-Meier (1958). To determine the implant

success, a wide scope of success criteria according to Albrektsson et al. (1986, Chart page 11) were used.

Wide scope of success criteria according to Albrektsson et al. (1986)

- Implant in situ
- Clear percussion sound
- Loosening degree 0–1
- Sulcus probing depths < 4 mm
- Absence of Bleeding On Probing
- Absence of clinical signs of infection
- Mean annual bone loss < 0.2 mm following the first year
- Subjective satisfaction with implant and dentures, no discomfort

Peri-implant pocket depths greater than 3 mm and up to a depth of 5 mm were recorded as peri-implant mucositis. Pocket depths greater than 5 mm met the criteria for periimplantitis and were therefore also a failure (Dvorak et al. 2011, Schmidlin et al. 2010, Zetterqvist et al. 2010, Gatti et al. 2008, Brägger et al. 2005).

Furthermore, failure was defined in the presence of the following examination findings: deep percussion sound associated with a loosening degree; positive Bleeding on Probing (BoP); clinical signs of infection such as redness, swelling, or pus discharge; a mean annual bone loss of more than 0.2 mm after the first postoperative year in combination with recession, pain, neuropathy, or dissatisfaction with the prosthodontic restoration as well as implant or abutment fractures.

## MIMI Nomenclature and Classification

In the end of 2021, the MIMI nomenclature of 2006 was subdivided by Nedjat to now 6 classes. In this study, cases were included in the following classes: MIMI 0 (immediate implantation), MIMI I (delayed implantation), MIMI II (horizontal distraction of narrow ridges without raising a mucoperiosteal flap), and MIMI VI (Internal Direct Sinus Lift, IDS). In 2011–2017, the French Matrix-bone augmentation material (Biomup, a collagen/beta-tricalcium phosphate) was used. Since 2017 Ethoss and/or the autologous, particulate dentin (Smart Grinder, Kometabio) has been used.

# MIMI® NOMENCLATURE

according to Dr. Armin Nedjat

- MIMI 0:** Immediate implantation  
(also with the Socket Shield (PET) technique and Smart Grinder procedure to process extracted teeth for grafting of autologous dentin)
- MIMI I:** Delayed implantation
- MIMI II:** Horizontal distraction
- MIMI III:** Vertical distraction
- MIMI IV:** Horizontal and vertical distraction
- MIMI V:** Indirect Sinus Lift (according to Summers)
- MIMI VI:** Internal Direct Sinus Lift (IDS according to Nedjat)

## Overview of patient gender distribution and number of placed and restored CHAMPIONS (R)Evolution implants with dentures

In this scientific study, 13,834 Champions (R)Evolution implants were placed.

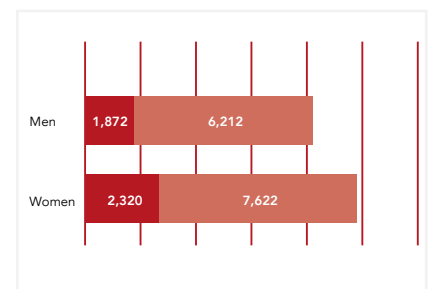
Of the patients 2,320 were women (55.34%) with a total of 7,622 implants and 1,872 men (44.66%) with 6,212 implants.

- Practice 1 (in Austria) placed a total of 3,117 CHAMPIONS (R)Evolution implants
- Practice 2 (in Germany) placed a total of 5,884 CHAMPIONS (R)Evolution implants

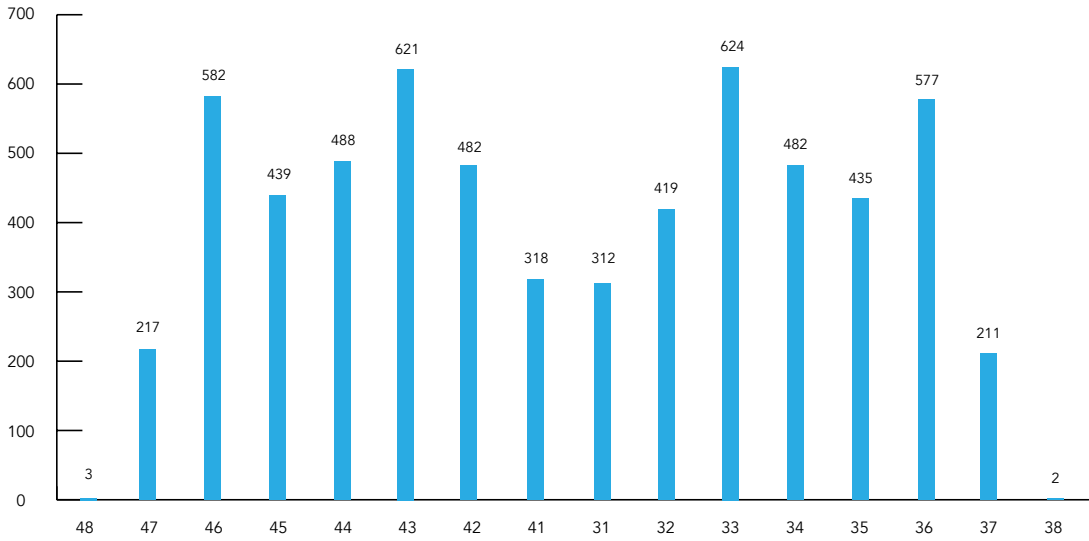
- Practice 3 (in France) placed a total of 2,406 CHAMPIONS (R)Evolution implants
- Practice 4 (in Poland) placed a total of 2,427 CHAMPIONS (R)Evolution implants

The mean age of the patients was 50.4 years at the time of implantation. Women received 3.26 implants on average and men 3.32 implants on average.

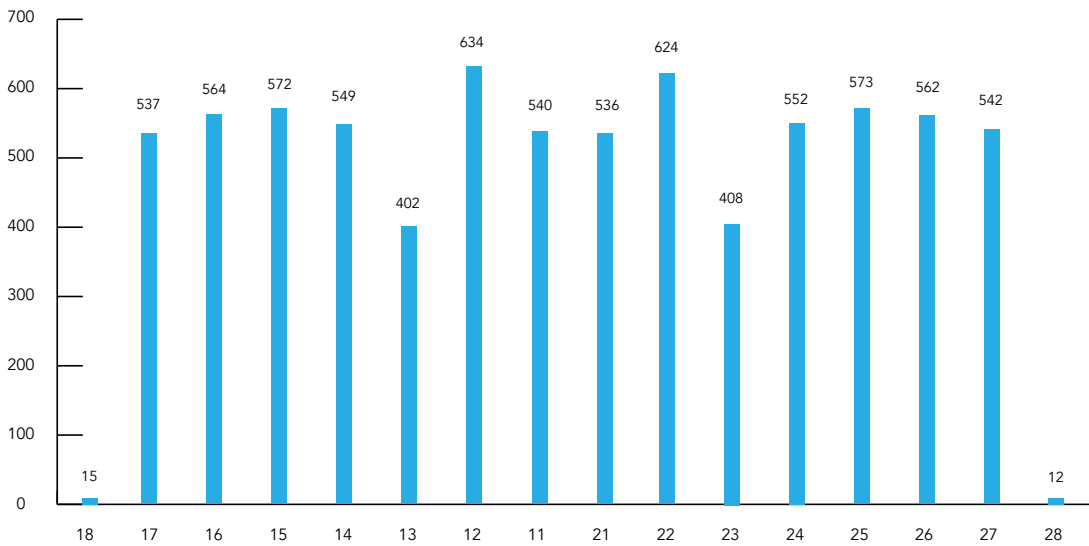
A total of 7,622 implants (55.1%) were placed in the maxilla. A total of 6,212 implants (44.9%) were placed in the mandible.

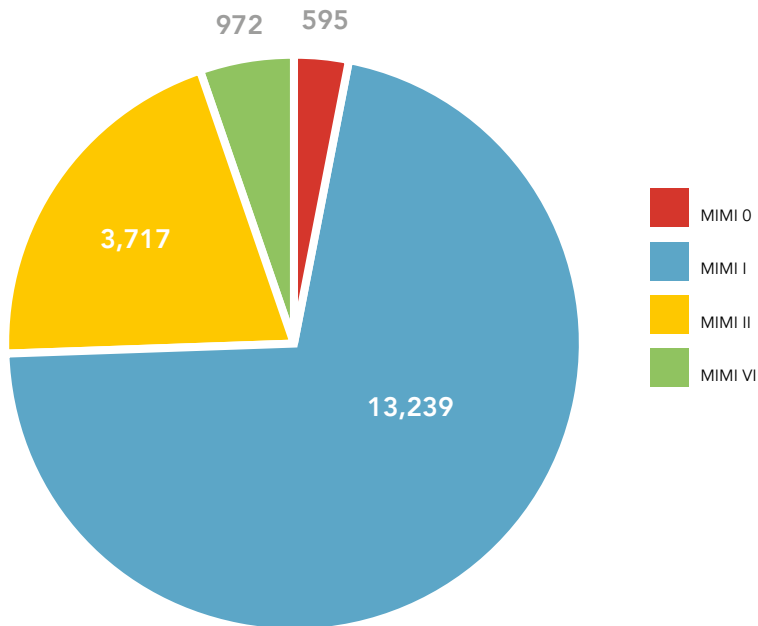
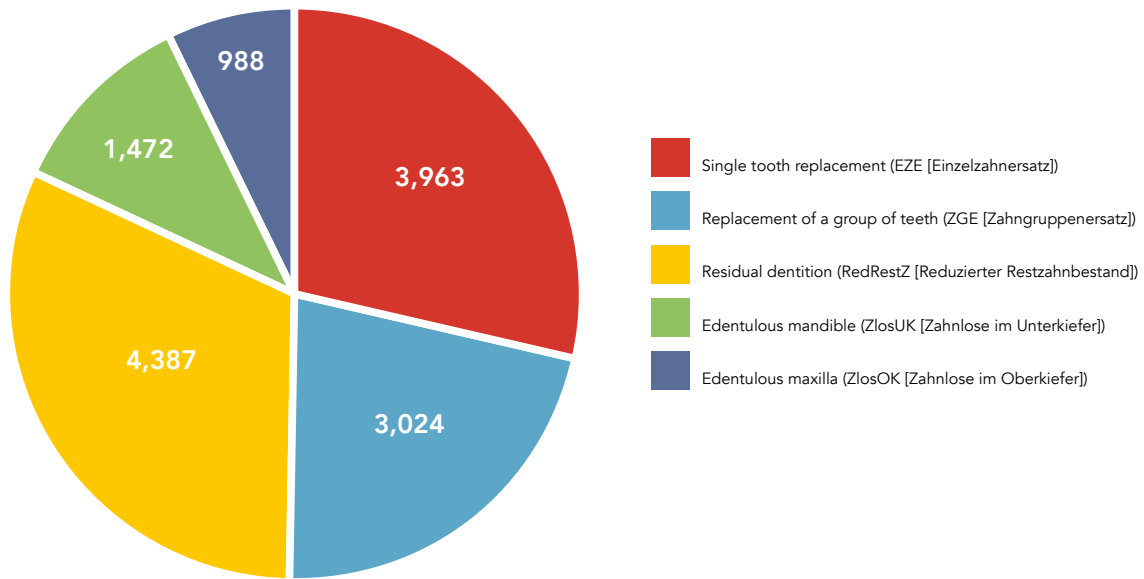


Distribution of 6,212 implants in the mandible according to the jaw region



Distribution of 7,622 implants in the maxilla according to the jaw region





### Distribution of all placed implants in the MIMI-classes

- Of a total of 13,834 implants there were 595 (4.3%) immediate implants (MIMI 0), i.e. tooth extraction and implantation in the same session.
- Most implants (95.70%) were delayed implants (MIMI I, from the 4<sup>th</sup> week post extraction).

- For delayed implantation, the horizontal, flapless distraction (MIMI II) was performed because of 28.08% preoperative soft tissue and hard tissue resorptions.
- Of 3,350 placed implants in the maxilla in the sites 15–17 and 25–27, 972 (28%) were performed with IDS (MIMI VI).

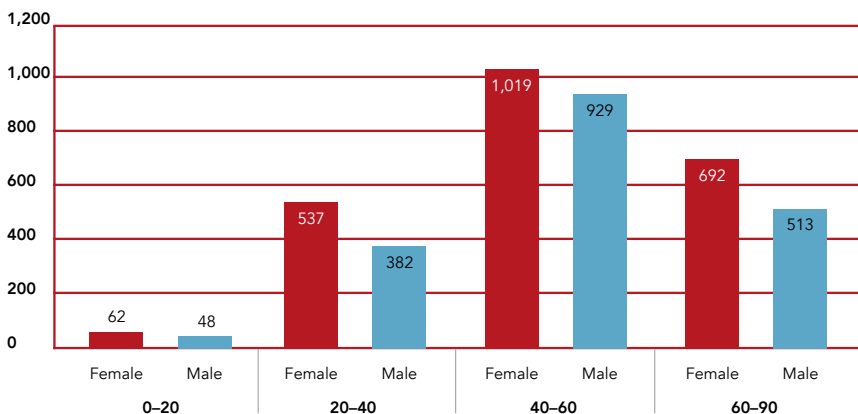
### Study population

Of the 4,192 patients who participated in the CIPC curriculum from 2011 to 2021 in the 4 clinics or under my supervision (Practice 2) and who received a total of 13,834 Champions (R)Evolution implants, 3,854 patients and 12,726 implants received follow-up care, and almost complete data was collected. This corresponds to a follow-up rate of 91.99%.

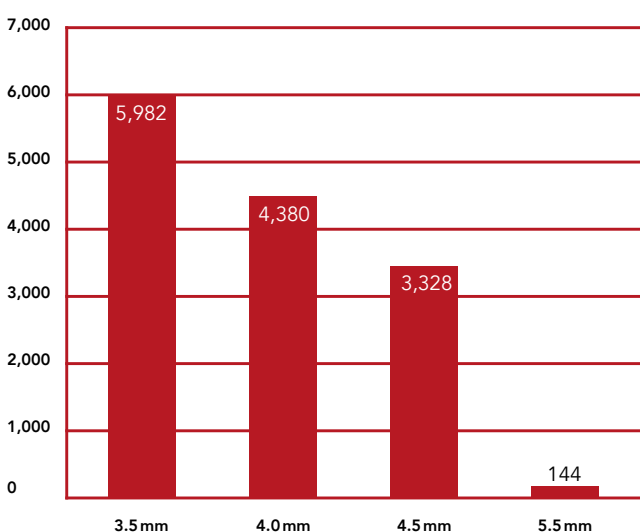
74 patients did not report back to their respective practices, 223 were unwilling to undergo a dental professional cleaning and follow-up visits due to their old age, serious illnesses, or personal reasons, 17 moved away unknown, and 24 died in the meantime.

The age groups were grouped together in 20-year increments (with the exception of the last age group: 60–90 years), starting with the group of 0–20 year-old patients. The youngest patient was 18 years old at the time of implant placement, the oldest was 86 years.

Age and gender distribution



Diameter of the placed and restored implants





**Detailed chart with implant diameter, divided according to respective lengths**

Influenced by “the guideline” of Prof. Dr. Jean-Pierre Bernard at the beginning of the study in 2011, short implants (< 12mm in length) with reduced implant diameters should be placed.

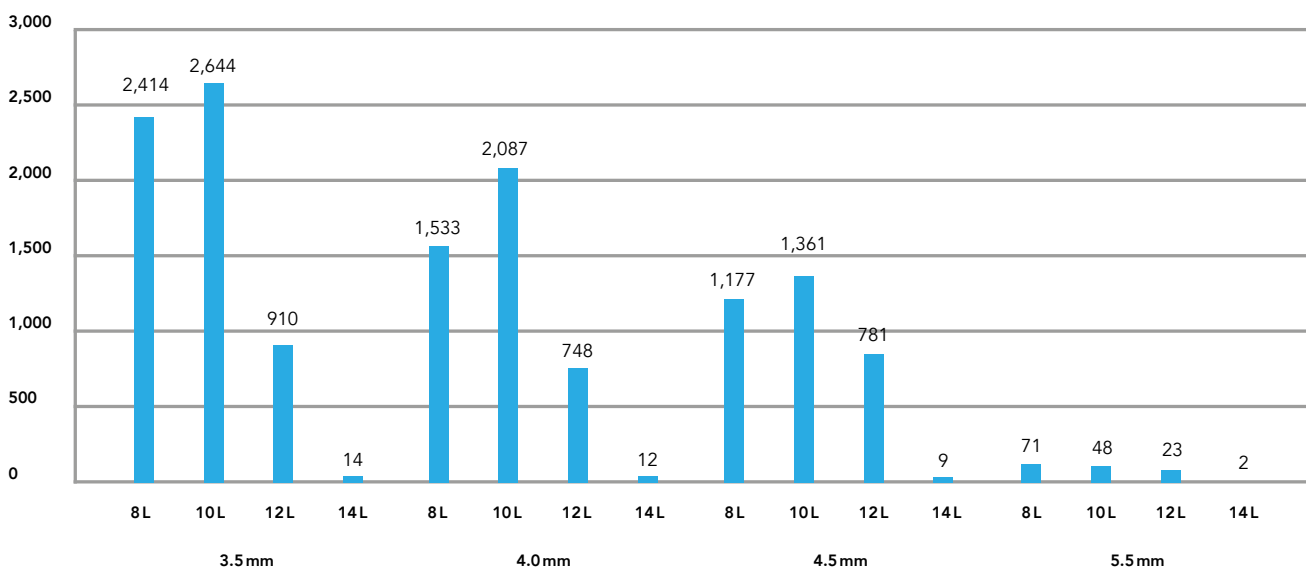
The other two guidelines for the treatment teams were that – if possible – the implants should be inserted 1–2 mm in the subcrestal position and that diameter-reduced (ø 3.5mm) implants with a primary stability between 20–40 Ncm should be placed.

**In addition to demographic data, the following parameters were collected:**

- Implant position
- MIMI-Nomenclature class 0–6
- Implant length and implant diameter
- Time of implantation
- Primary stability in Ncm
- Equicrestal, subcrestal, and supra-crestal insertion
- Augmentation: yes/no?
- Indication class [single-tooth dentures (Einzelzahnersatz [EZE]), replacement of a group of teeth (Zahngruppenersatz [ZGE]), residual dentition (reduziertes Restgebiss [RRG]), edentulous maxilla (zahnloser Oberkiefer [ZOK]), edentulous mandible (zahnloser Unterkiefer [ZUK])

- Day of the restoration with the superstructure
- Type of superstructure
- Bone loss and tissue loss in mm
- Complications
- Day of the implant failure
- Reason for loss
- Recovery of stability after “tightening” in case of loosening of the implant within the first 4 weeks after surgery (“Erni-Test”)
- Day of the last check-up

The data material was recorded in charts and statistically processed concerning individual questions.



### Requirements for practices for studies of implantation and dentures:

- After insertion, each implant should have an attached gingiva of a minimum width of 1 mm in the vestibular position.
- Each implant with the Shuttle should end in a supragingival position at a maximum of 1 mm.<sup>X1</sup>
- Each implant should achieve minimum primary stability at 20 Ncm.
- In the D1/ D2 bone there should be a "crestal relief" of 1–2mm.
- As a rule implants can be immediately restored/immediately loaded if 4 implants/teeth are attached/splinted with a passively fitted temporary denture or the final superstructure.
- If possible, short implants of 8 mm and 10 mm should be placed.
- The primary goal is to place an im-

- plant diameter of only 3.5 mm and of only 4.0 mm for single molar implants
- Until you fit the superstructure, you should wait 3 months for an immediate implantation (MIMI 0) and MIMI-II (horizontal distraction), 4 months for MIMI-VI (Internal Direct Sinus Lift), and 2 months for a "classical" delayed implantation (MIMI I).
- Crowns should be fitted with phosphate cement or Relyx Unicem (3M Espe) and not temporarily.
- The Abutments should reach a final torque of 30Ncm. In the posterior maxilla (D3/D4 bone) the final torque should be 25Ncm.

X1: From January 2018 this requirement was corrected so that it was now possible to place a Shuttle with an equigingival closure or the Hybrid Screws Georgi with a gingival height

of GH 1.5 (golden) and GH 2.5 (pink), which have then been introduced, in addition to the already existing Surgical Cover Screws with GH 0.5.

A Hybrid Screw Georgi should preferably be closed at 1–2 mm in a subgingival position. The Hybrid Screws Georgi are available in gingival heights of GH 0.5 (gray), 1.5 (golden), and 2.5 (pink).

Two-piece implant systems have therefore been developed to strictly avoid lateral micromovements within the first 8 weeks post-surgery. During this time, peri-implant bone is broken down, remodeled, and modeled. If, for example, a CHAMPIONS (R)Evolution implant is placed at a mucosal thickness of 2 mm with its 3.5 mm-high Shuttle at "bone level", its osseointegration is jeopardized in the "healing

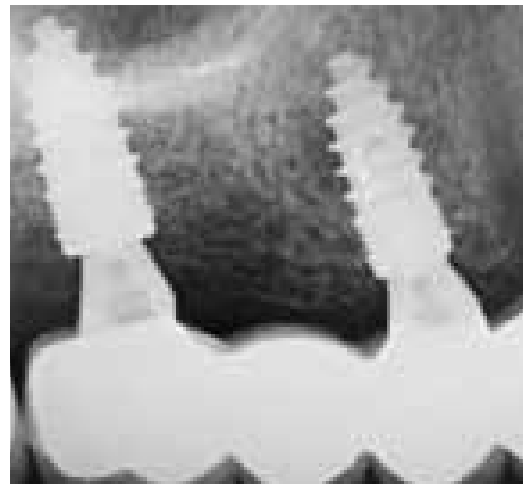


phase" since only the slightest movements can badly affect the implant osseointegration. External factors such as tongue movements on the implant would thus be able to endanger the osseous implant integration. In these cases, after taking implant X-rays-still with the Shuttle – you should remove the screw and the Shuttle in order to replace the Shuttle with one of the three "Hybrid Screws Georgi". In this way, there are no micromovements of the CHAMPIONS (R)Evolution implant due to eating behavior or a removable denture.

If you leave the 3.5mm-high Shuttle sterile on the implant in case of a gingival height of e.g. 2mm, subcrestal implant placement is recommended. Then the Shuttle also closes at "tissue level" or even in a

slight subgingival position so that no movements and forces on the implant can occur during the implant healing phase. Subcrestal implantation is very possible with the CHAMPIONS (R)Evolution system as due to the double cone with integrated Hexadapter, there is no micro-gap that is vulnerable to bacteria. Due to this subcrestal implantation procedure, the exchange of the Shuttle for the Hybrid Screw Georgi is not necessary.

Another advantage of the subcrestal implantation is that at crestal bone level there are no great forces so that a crater-shaped bone loss during the healing phase can be avoided.



## Navigation

There are 3 navigation aids:

CNIP-Navigation: actually, CNIP, the "Cortical Navigated Implantation Procedure", is a natural procedure. CNIP is performed using Conical Triangular Drills at low speed (about 50–70 RPM in the spongy bone) and guided through the cortical structures of the bone (250 RPM), which ensures that the drills always stay in the spongy bone without perforating cortical structures. Using

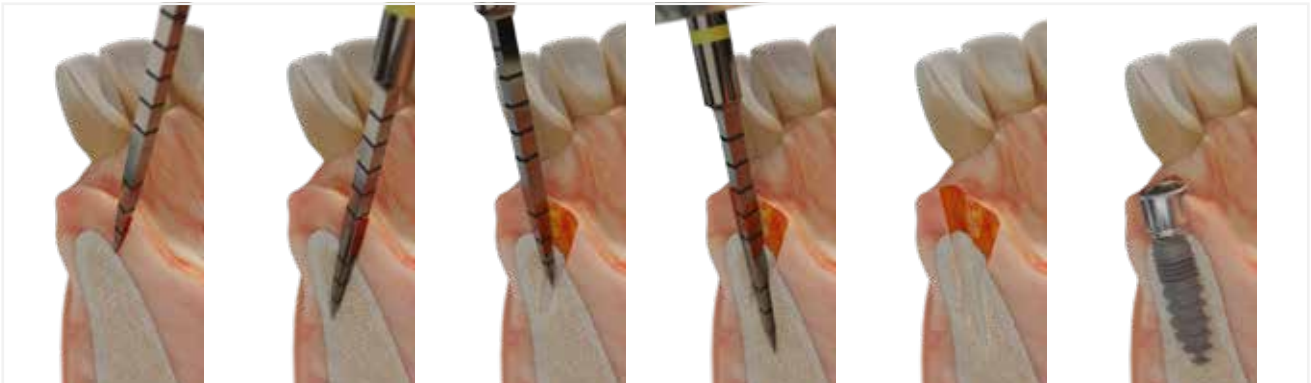
CNIP, cortical bone structures in the buccal and oral positions serve as natural navigation aids.

With a correct procedure and by performing a BCC (Bone Cavity Check) with a sufficiently long, thin, flexible metal probe after the first and last drilling, you can successfully and safely prepare the bone for the flapless implant placement.

The "Chetry" Drill Stop Sleeves for the yellow pilot drill are another aid for

finding the correct distance between several implants next to each other. With the "Chetrys"  $\varnothing$  5.0,  $\varnothing$  7.0 and  $\varnothing$  9.0mm, the ideal implant spacing is automatically obtained by leaving the yellow drills together with the Drill Stop Sleeve inserted in the drilling position.

For the user CHAMPIONS Guides serve as a navigation aid for the optimized prosthodontic position, allowing for preoperative and intraoperative individual shaping for the patient.





### Implantation in the dense D1/D2 bone

For a "Bone-Level" implantation in dense D1/D2 bone, crestal bone is relieved:

- For a 3.5mm-diameter implant, the 4.0mm-diameter drill is finally used
- For a 4.0mm-diameter implant, the 4.5mm-diameter drill is finally used in the crestal position



QR Code: Dr. Nedjat explains "Bone Level", "Tissue Level", subcrestal implantation "crestal relief" and "platform switching"



Surgery and prosthodontic restoration of a CHAMPIONS (R)Evolution implant in site 46



## Implantation in the D3/D4 bone

In the low density bone (D3+D4) and for immediate implantation, the so-called "osseous metamorphosis" (OM) occurs. Using your hands, so "digitally" (manually and with the Torque Wrench Insertion Aid), the first two Conical Triangular Drills are used making awl-like movements. Then, only the CHAMPIONS Condensers are used.

Condensers are bone condensing instruments. The Condensers are located in the 2<sup>nd</sup> row of the CHAMPIONS system: if you achieve primary stability at a torque of about 20Ncm (hand-tightened) with the Ø 4.3mm blue Condenser in the low density bone, place a 4.5mm-diameter implant. Condensers also allow you for intraoperative determination of the length of the required implant.

- The anatomy (e.g. by means of a preoperative cone beam) does not determine which implant diameter is to be used. Rather, the diameter of the implant is determined by the bone density; Condensers allow for intraoperative determination of the bone density in the low density bone or when performing immediate implantation.



## Which implant Ø do you place in low density bone?

### First intention:

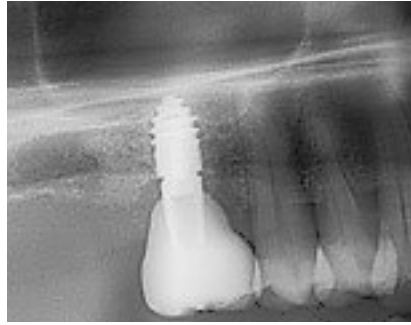
Ø 3.5 mm when primary stability at 30/40 Ncm is achieved

Exception: single molar  
→ Ø 4.0 mm

Condenser stability at 20 Ncm

Implant diameter

	3.3	→	3.5	
	3.8	→	4.0	
	4.3	→	4.5	
	5.3	→	5.5	



**Case report 6:** For immediate implantation for maxillary first molars, a tri-furcation is performed, which is also recommended from a prosthodontic point of view. Immediate implantation in the mandibular molar region can be successfully performed by means of Condensers in a few minutes. Spongy bone can be modeled very well so that a 4.0mm-diameter implant can achieve primary stability even in a thin, 1.5mm-wide septum.



Film: Immediate implantation in the molar region

### Scientifically prepared data of 2011–2021

Observations included patients who underwent implant therapy in 4 Implantology dental offices in Europe (one in Austria, one in Germany, one in France, one in Poland) between July 2011 and September 2021. Observations (Practice 2) also included all CHAMPIONS (R)Evolution cases that have been treated since February 2017 by Implantology novices of the one-year CIPC Implantology & Implant Prosthodontics Curriculum by two experienced supervisors across Germany. This intensive, comprehen-

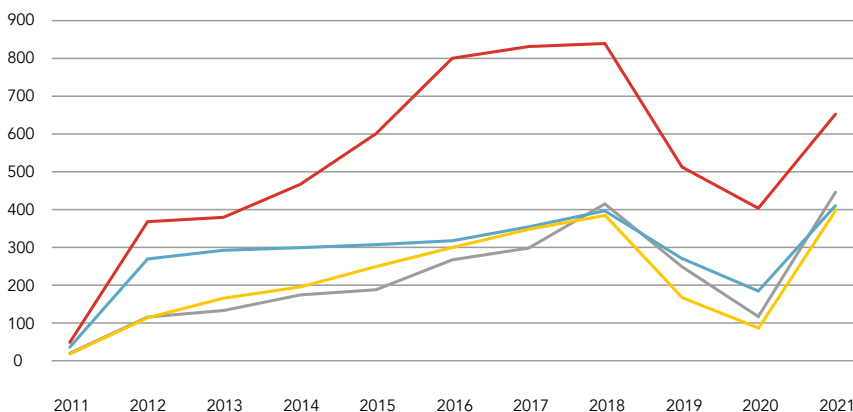
sive training concept is based on the SIAO/CITC concept of Prof. Dr. Jean-Pierre Bernard (University of Geneva), who advised his students in their respective private dental offices within the framework of the Clinical Implant Training Concept in the 90s.

### Iatrogenic influence factors: learning curves of the practitioners

Practitioner teams needed some time to adjust when introducing a new implant system in their practice. Placing this implant system has become routine after treating about 10 patients. After the first surgeries in 2011, the

procedure became more routine in the following years and reached a preliminary maximum after 7 years with more than 800 implants (Practice 2) per year. Due to lock-down measures and uncertainties in individual countries because of the Corona pandemic, all practices were not able to continue implanting to the same extent in 2019 and 2020 as they had been able in the years before, but then they increased again implantations significantly in the year 2021.

Implants/year in the individual practices from 2011 to 2021



**Overview of inserted implant lengths and diameters as well as other parameters in the involved practices**

In the frame of this scientific study, 13,834 implants were placed and re-stored with dentures in 4,192 patients. The chart below shows an overview of the placed CHAMPIONS (R)Evolution implants:

A%V Augm.: Number and % of failures with simultaneous augmentation

PSt V: Primary stability in Ncm for failures on average

A%V GH > 0mm: Number and % of failures when during the "healing phase", the Shuttle or the Georgi was placed with a gingival height of more than 0 mm

II ∑&%: Immediate implantation, number and % of implant types

Failure II ∑&%: Quantity of immediate implant failures and % of implant types

Implant length / Ø in mm	Practice 1		Practice 2		Practice 3		Practice 4		Total		Number of failures and %	A % V Augm.	PSt V	A % V > GH 0 mm	II ∑ & %	Failures II ∑ & %
	N	%	N	%	N	%	N	%	N	%						
CHAMPIONS (R)Evolution 8 – 3.5	582	18.67	1,135	19.42	272	11.31	425	17.51	2,414	17.45	84 3.48 %	5 5.95 %	< & = 20 Ncm	62 73.81 %	2 0.08 %	0
CHAMPIONS (R)Evolution 8 – 4.0	118	3.79	851	14.56	187	7.77	377	15.53	1,533	11.08	52 3.39 %	3 5.77 %	< & = 20 Ncm	38 73.08 %	12 0.78 %	0
CHAMPIONS (R)Evolution 8 – 4.5	84	2.69	486	8.32	252	10.47	355	14.63	1,177	8.51	42 3.57 %	13 30.95 %	< & = 20 Ncm	38 90.48 %	56 4.76 %	2 3.57 %
CHAMPIONS (R)Evolution 8 – 5.5	12	0.38	52	0.89	3	0.12	4	0.16	71	0.51	2 6.39 %	2 100 %	< & = 20 Ncm	2 100 %	4 5.63 %	1 25 %
CHAMPIONS (R)Evolution 10 – 3.5	876	28.10	945	16.17	386	16.04	437	18.01	2,644	19.11	91 3.44 %	4 4.39 %	< & = 20 Ncm	78 85.71 %	21 0.79 %	1 4.76 %
CHAMPIONS (R)Evolution 10 – 4.0	547	17.55	734	12.56	404	16.79	402	16.56	2,087	15.09	71 3.40 %	6 8.45 %	< & = 20 Ncm	62 87.32 %	13 0.62 %	0
CHAMPIONS (R)Evolution 10 – 4.5	391	12.54	342	5.85	276	11.47	352	14.50	1,361	9.84	47 3.45 %	8 17.02 %	< & = 20 Ncm	41 87.23 %	124 9.11 %	4 3.22 %
CHAMPIONS (R)Evolution 10 – 5.5	2	0.06	31	0.53	6	0.25	9	0.37	48	0.35	1 2.08 %	1 100 %	< & = 20 Ncm	1 100 %	0	0
CHAMPIONS (R)Evolution 12 – 3.5	274	8.79	422	7.22	188	7.81	26	1.07	910	6.58	34 3.74 %	2 5.88 %	< & = 20 Ncm	30 88.24 %	112 12.31 %	2 1.79 %
CHAMPIONS (R)Evolution 12 – 4.0	185	5.94	327	5.60	221	9.19	15	0.62	748	5.41	29 3.88 %	2 6.90 %	< & = 20 Ncm	18 62.07 %	66 8.82 %	1 1.52 %
CHAMPIONS (R)Evolution 12 – 4.5	32	1.03	534	9.14	202	8.40	13	0.54	781	5.65	28 3.59 %	5 17.85 %	> & = 40 Ncm	20 71.43 %	145 18.57 %	2 1.38 %
CHAMPIONS (R)Evolution 12 – 5.5	1	0.03	16	0.27	4	0.17	2	0.08	23	0.17	1 4.35 %	1 100 %	> & = 40 Ncm	1 100 %	15 65.22 %	0
CHAMPIONS (R)Evolution 14 – 3.5	6	0.09	4	0.09	2	0.08	2	0.08	14	0.10	0 0 %	0	-	-	9 64.29 %	0
CHAMPIONS (R)Evolution 14 – 4.0	5	0.09	2	0.03	2	0.08	3	0.12	12	0.09	1 8.33 %	0	-	-	7 58.33 %	0
CHAMPIONS (R)Evolution 14 – 4.5	2	0.06	2	0.03	1	0.04	4	0.16	9	0.07	1 11.11 %	0	-	-	7 77.78 %	0
CHAMPIONS (R)Evolution 14 – 5.5	0	0	1	0.02	0	0	1	0.04	2	0.01	0 0 %	0	-	-	2 100 %	0



## Statistical methods

Statistical calculations performed using SPSS 11.0.0 (IBM, Armonk, NY, USA) and SAS Version 9.2 (SAS Institute Inc., Cary, NC, USA). In the context of the available analyses, the following characteristic values were given, depending on the following values:

- For frequency data, these were absolute and/or relative frequencies (percentage values).
- For metric data there were the arithmetic means, as a measure for variability, the standard deviation, the minimum and maximum, the number of cases, and percentiles.
- Survival rates have been presented as a Kaplan-Meier-curve.
- Individual subgroup significances were investigated using log-rank or chi-square test statistics. The p-values and test statistics were reported.
- In case of statistically significant group differences, the estimated difference (%) as well as its 95% confidence interval are reported.

## Results

### Implant-related survival rate and survival probability

The 1-year survival probability of all implants was 98.89% and decreased to 97.6% after 5 years and 96.5% after 10 years.

The observed overall failure rate (failure rate) of 3.50% improved to 2.40% when the CHAMPIONS (R)Evolution implants were placed 1–2 mm in the subcrestal position.

### Reason for implant failures

Altogether, 484 of 13,834 implant failed during the 10-year observation time, corresponding to 3.50%. For the individual practices the values were 2.9% (Practice 1), 4.2% (Practice 2), 3.2% (Practice 3), and 3.7% (Practice 4).

Implant failures were predominantly related to the supragingival position of the Shuttle (thanks to the MIMI methodology not colonized by bacteria, i. e. non-inflammatory) within the first 8 weeks after surgery. Acute and chronic peri-implantitis as well as classic peri-implantitis affecting soft tissue and bone contributed only very little to implant failures. The guidelines or statements of the leading scientific societies usually refer to the late form of periimplantitis with mucositis and ostitis, which we refer to here as periimplantitis chronica.

The early forms of inflammations, which also lead to loss, are given less consideration. These are the acute osteomyelitis or its localized form, which immediately force you

to remove the implant (periimplantitis totalis acuta) or the insidiously progressing form, which the patient does not notice, and the practitioner is only surprised to discover that there has been very little or no osseointegration in the prosthodontic phase, with the consequence that the implant has a degree of loosening II (periimplantitis totalis chronica). From a patho-histological point of view, these are also inflammations, although they are completely different from the peri-implantitis that occurs later with mucositis and progressive bone loss around the implants after initial osseointegration.

Most implant failures occurred in the first months during the healing period of the implants: In the first 2-4 months post implantation, 436 implants (90.08% of failures) failed, or 3.15% of all placed implants failed.

By the end of the first year after fitting the denture, 27 (5.58% of the failures) or 0.2% of the remained implants with superstructure failed, another 13 implants (2.69% of the failures) by the end of the second year or 0.1% of the remained implants were lost, up to 5

years another 5 (1.03% of the failures) or 0.04% of the remained implants were lost, up to 10 years 3 (0.62%) or 0.02% of the remained implants were lost.

Only 3 implants failed because of implant fracture: a CHAMPIONS (R)Evolution implant with a length of 14 mm and a diameter of 3.5 mm in 46, an implant (12 x 3.5 mm) in 13 (bridge sites 13–16), and an implant (10 x 3.5 mm) in 36 (as splinted crowns 36+37).

Since the Hybrid Screws Georgi were introduced in the year 2017, a progressive decrease in the failure rate in the first 2–4 months of healing time was observed. When the Shuttle sometimes protruded more than 0.5 mm from the gingiva during equicrestal implantation, it therefore led to lateral micromovements of the implant (e.g. through a temporary removable denture). By replacing the 3.5 mm-high Shuttle with a Surgical Cover Screw Georgi with a gingival height (GH) of 0.5 mm, a Hybrid Screw Georgi with a GH of 1.5 mm (golden), or the one with a GH of 2.5 mm (pink), complete non-osseointegration or insufficient osseointegration could be significantly

Reason for explantation	N (Implants)	% (Failures)
Periimplantitis chronica	45	9.30
Periimplantitis totalis chronica	47	9.71
Periimplantitis totalis acuta	8	1.65
Osteolysis/ Residual necrosis in case of immediate implantation	12	2.48
Lateral forces exerted on the implant during the healing time (Shuttle > 1 mm in supragingival position), no signs of inflammation, no pain	324	66.94
Implant fracture	3	0.62
Osteolysis in case of augmentation with foreign bone replacement graft	1	0.21
Aversion, explantation on the patient's demand	2	0.41
Paresthesia	1	0.21
(Grade 3 & 4) Ti-O <sub>2</sub> intolerance	41	8.47
Total	484	100.0

reduced in these cases during the healing phase.

As long as the CHAMPIONS (R)Evolution system was placed at least 1–2 mm in a subcrestal position (with a gingival height of e.g. 2 mm), there were almost no osseointegration failures of the implants during the healing phase of 2–4 months.

The dogma “Implants should be as long and wide in diameter as possible”, which was propagated at the beginning of the history of Implantology, has been impressively refuted (also by preliminary investigations by Prof. Dr. Jean-Pierre Bernard, University of Geneva/Switzerland).

- It has been preferable to place an 8 mm long implant, correctly placed in the ideal prosthodontic position (the old rule of always placing a 4.0 mm diameter implant for a single molar has been observed) rather than a longer implant placed as close as possible to the inferior alveolar nerve. There were no statistical differences in implant failures related to implant lengths of 8, 10, 12 and 14 mm for the same diameters and

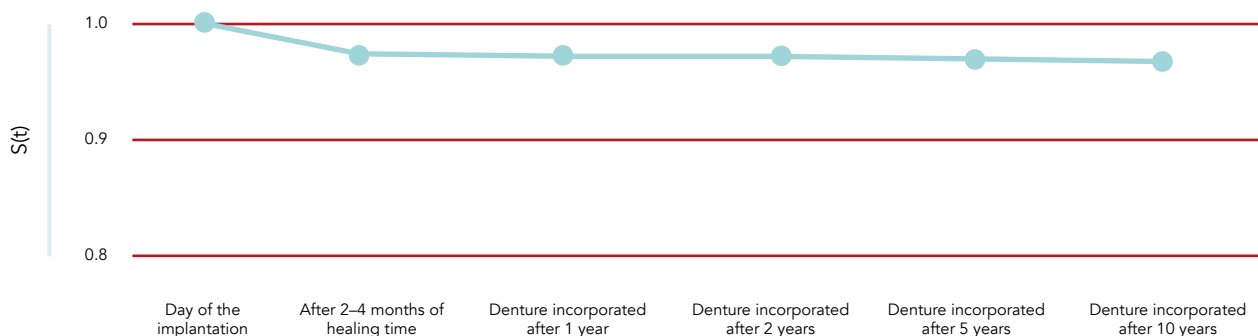
maintenance of adequate primary stability.

Surprisingly, there were no increased complications or failures in case of an immediate implantation (extraction and implantation only in one treatment session → MIMI 0) compared to a delayed implantation (MIMI I): Of a total of 595 immediate implants (4.3% of all implantations), only 13 implants failed. This corresponds to a failure rate of only 2.18% compared to a total of 3.5% (or 2.4% “corrected”). This is probably due to the good surgical protocol with the help of the Condensers, the periosteum preserving surgery concept, and the bone condensing design of the CHAMPIONS (R)Evolution system:

- Bone-preserving cavity preparation using Conical Triangular Drills and Condensers.
- Palatal/ lingual axis inclination to protect the buccal bone wall with fully intact periosteum
- Achieving at least 40 Ncm primary stability only through the spongy bone
- No buccal pressure on the buccal bone lamella

MIMI, the Minimally Invasive Implantation Method, has become commonplace in the surgical and prosthodontic dental practice, meeting patients’ needs. Dental Implantology has been revolutionized by the MIMI procedure for over 25 years: mucogingival flap detachment, including periosteal detachments, and reopenings in the prosthodontic phase with multiple manipulations inside the implant are not only time-consuming and unnecessary, but also the main reasons for iatrogenic peri-implantitis and subsequent complications. The main advantage of MIMI, however, is that it is time-saving and efficient so that dental implantations can be performed effortlessly every day (and also “spontaneously” such as an immediate implantation). Thus, Implantology has become a completely demystified regular discipline in a regular dental office.

S(t) for ALL implants



## Summary

The aim of this clinical, retrospective study was to investigate the long-term survival rate and success rate of the CHAMPIONS (R)Evolution implant system and to calculate the statistical long-term survival probability and the success probability as well as to compare them with each other and with data of previous studies. This should help both the practitioner and the patient in making treatment decisions by ensuring the greatest possible safety regarding the predictability of the long-term success of this implant system.

In this study, 4,192 patients received a total of 13,834 CHAMPIONS (R)Evolution implants within 10 years, from September 2011 to September 2021.

Data were collected concerning age, gender, date of the implant placement, and the clinical examination, augmentations performed with the augmentation material used, time between tooth loss and implant placement, the subsequent prosthodontic restoration and the presence of risk factors for implant success.

The percussion sound of the implant, the loosening degree, the sulcus probing depths, and recessions in 4 areas were clinically studied. Bleeding On Probing as well as signs of inflammation of the peri-implant mucous membranes were documented.

Regular X-rays were taken to evaluate bone loss.

Patient satisfaction and the psychosocial aspect of implant care were assessed with a questionnaire. The data obtained were statistically evaluated using Cox regression analysis, and survival probabilities were determined according to the Kaplan-Meier method.

After an average of 6 years, 96.5% of the implants inserted in the minimally invasive MIMI procedure were still functioning, and the success rate according to the extended criteria of Albrektsson was 92.5%.

The main cause of implant failure was failed healing in 67% of the cases in the first 12 weeks post implantation, which, however, could be statistically reduced

to only 20% by the "Erni test" ("re-tightening of the implant" in the 3<sup>rd</sup>/4<sup>th</sup> week from 2011–2016) and by the introduction of the two additional Hybrid Screws Georgi (GH 1.5 – GH 2.5) in 2017 for equigingival healing.

There were 31.82% of inflammatory processes leading to implant failure.

Thus, the minimally invasive implantation method (MIMI) with its periosteum-preserving procedure is a very efficient surgical technique, which fully ensures bone nourishment even for years.

Nicotine abuse and chronic periodontitis had a statistically significant influence on the survival probability.

Augmentations in the course of MIMI II (horizontal distraction) and MIMI VI (Internal Direct Sinus Lift), the implant length, and the implant diameter had no influence on implant failure, as long as the given parameters for implant placement were taken into account (primary stability and protection against lateral micromovements).

The CHAMPIONS (R)Evolution implant system can be considered as safe therapy alternative to bridges or conventionally retained dentures.

It can also be safely assumed that the failure rates are better with limited indications than with implantation in the full range of possibilities. In addition, the anamnestic patient selection seems to play a role, also with the MIMI procedure, in particular with immediate implantation (MIMI 0) and MIMI II/MMI VI.

Thus, it also seems justified to state that inexperienced people can achieve good results. The failure rates at the beginning of implantological activity can be almost equally low with the CHAMPIONS (R)Evolution with previous (online and practical) training. Once you get familiar with the field of Implantology and the MIMI procedure, the failure rate will be as low as for experienced users.

A difference between the indication classes has not been observed.

Also, a horizontal distraction (MIMI II) or IDS (Internal Direct Sinus Lift) did not play a role in the implant failure, although there was still a trend in favor of implants in the local bone.

Implants were equally successfully placed and restored with dentures in the maxilla or in the mandible ( $p = 0.0001$ ).

There is a 96.5% probability that an implant is still incorporated after 10 years if the implant is placed in an equicrestal position. If necessary, the Shuttle must be exchanged for one of the 3 Hybrid Screws Georgi in order to close it at least in an equigingival position, if not then even in a slightly subgingival position. With this procedure, the "5-year study" of 2016 showed a survival probability of 97.6%.

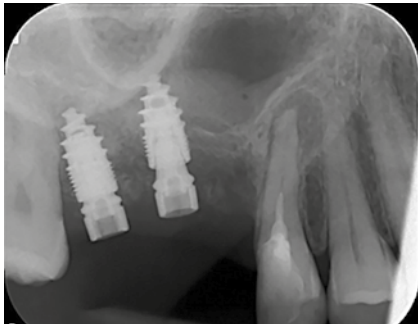
The innovation of the CHAMPIONS (R)Evolution design with crestal microthread and the inner double cone of 9.5°, the surface finishing through Al<sub>2</sub>O<sub>3</sub> radiation and triple etching of the implant surfaces, which has remained unchanged since 2011, the so far unique innovation of the Shuttle, which is attached to the actual implant with a Screw with 10Ncm, and last but not least the cost-effectiveness for end customers are certainly reasons for the success of the system in Europe.

The sustainability of CHAMPIONS (R)Evolution has been scientifically proven in this 10-year study.

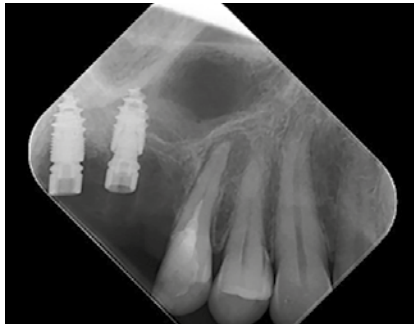
Meanwhile, the CHAMPIONS (R)Evolution system is a tried and tested, fully developed implant system, which is completely convincing in surgery (also thanks to the non-traumatic and patient-friendly minimally invasive insertion), and in Prosthodontics (e.g. only one prosthetic platform for all implant diameters) and in its optimized workflow.

X-rays of case series with the CHAMPIONS (R)Evolution implant system

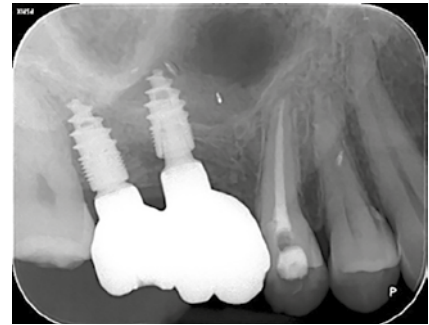
Case 1: Implantation sites 16 + 17 (with sinus lift 16)



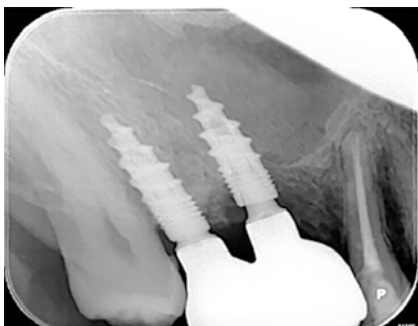
12.09.2011



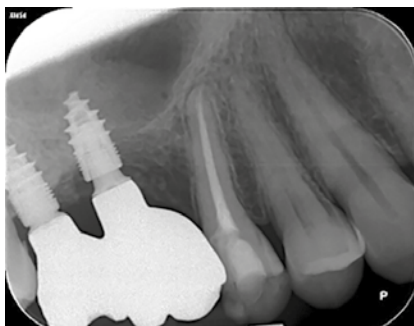
14.02.2012



09.03.2016

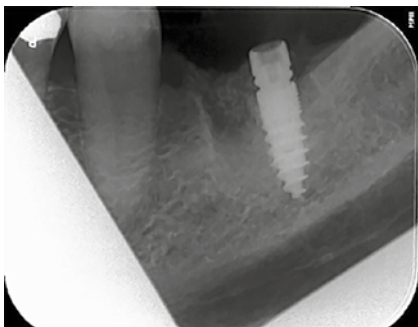


07.03.2018

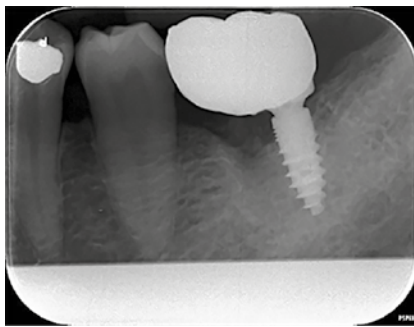


18.10.2021

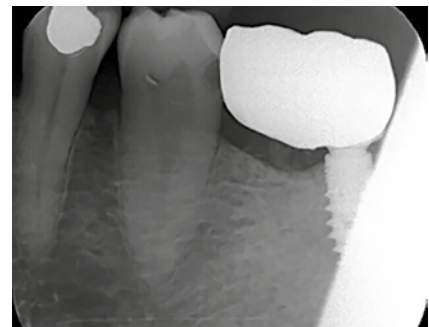
Case 2: Implantation site 36



12.06.2014

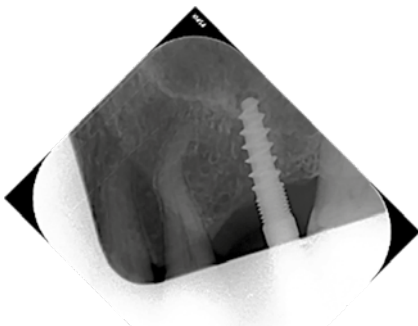


10.10.2014

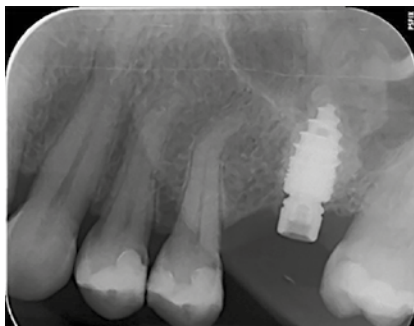


05.03.2021

Case 3: Implantation in site 26 with simultaneous IDS sinus lift surgery



28.02.2014

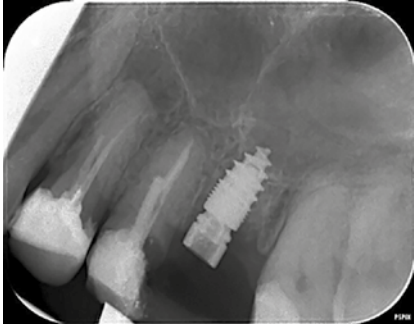


28.02.2014

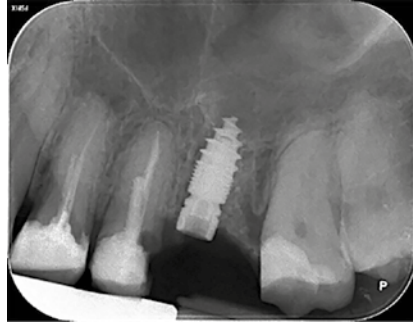


22.03.2021

Case 4: immediate implantation in site 26 with simultaneous augmentation



17.10.2011



17.10.2011

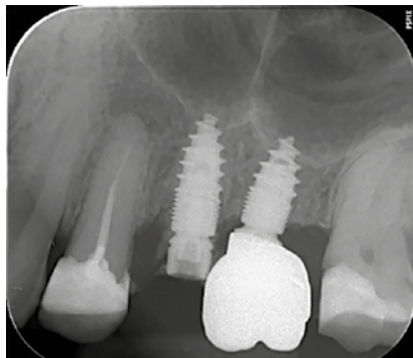


24.01.2013

Case 5: immediate implantation site 25 (same patient as Case 4)



13.03.2015

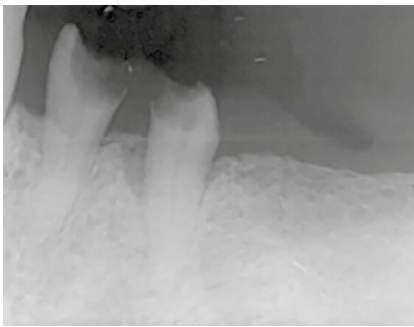


13.03.2015

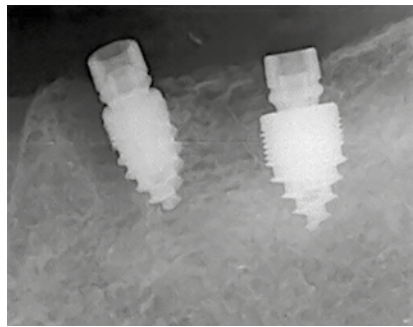


28.08.2018

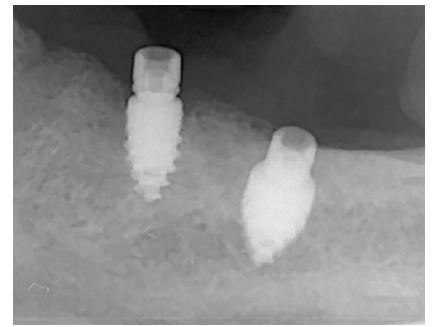
Case 6: immediate implantation sites 34 + 36



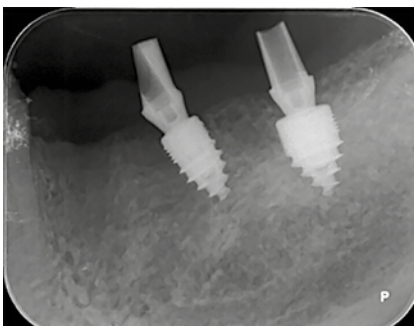
12.12.2011



12.12.2011

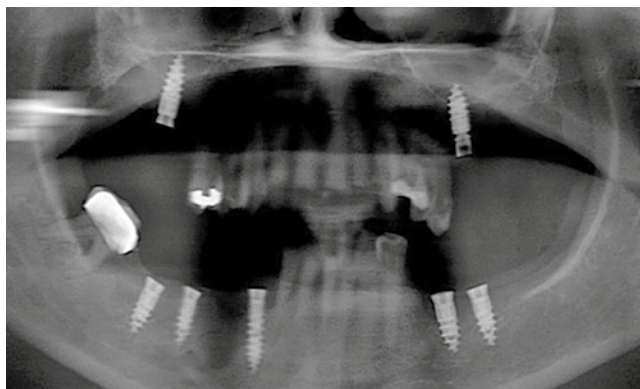


03.04.2012



14.12.2021

Case 7: immediate implantation 17 +  
27 + 36 + 35 + 44-47

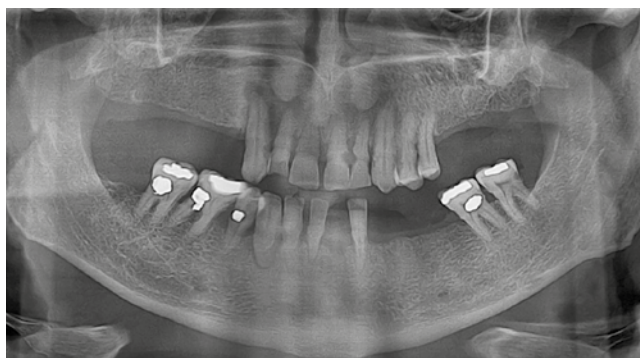


12.05.2015



09.08.2021

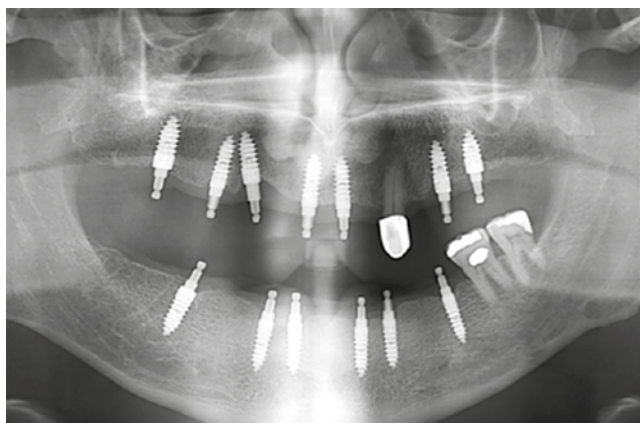
Case 8: Implantations 17, 15, 13, 11,  
21, 25, 25, 35, 33, 32, 42, 43, 46



21.11.2011

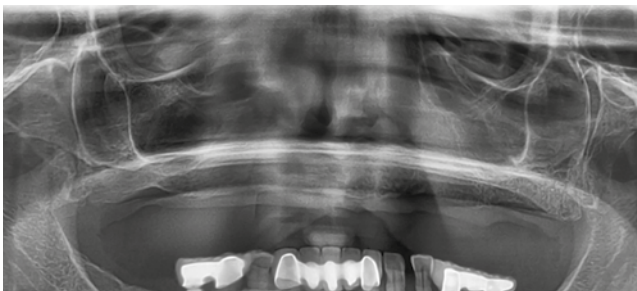


03.02.2012

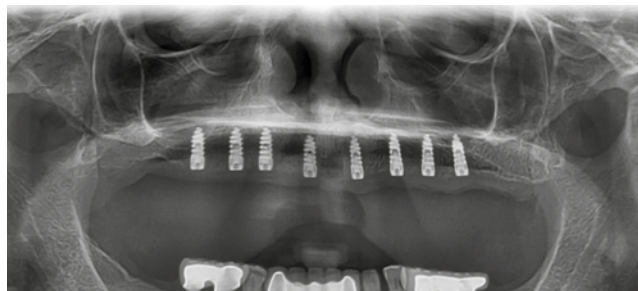


27.08.2021

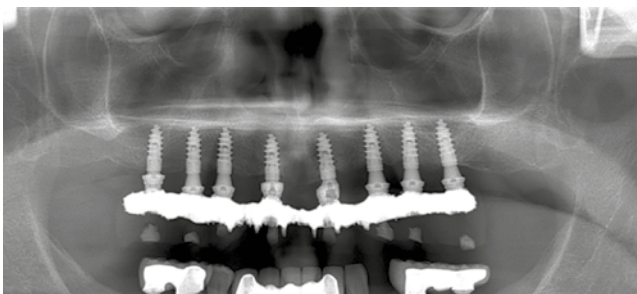
Case 9: late implantation in the maxilla in sites 15-25



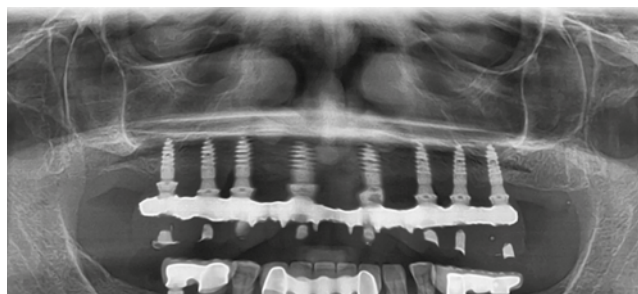
06.01.2011



23.09.2011

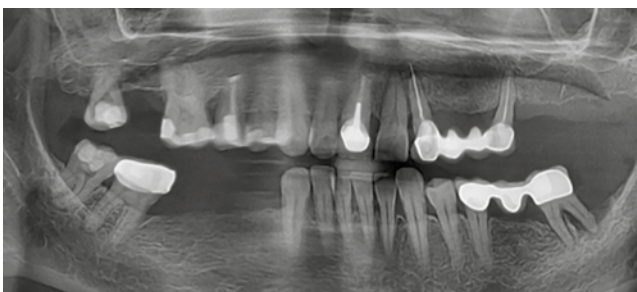


17.08.2012

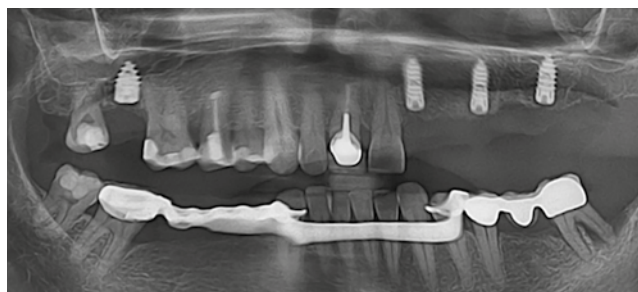


24.04.2020

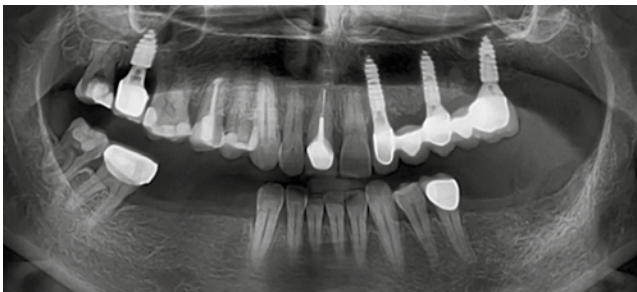
Case 10: Implantations sites 17, 22, 24 + 26 as well as 36, 44 + 46



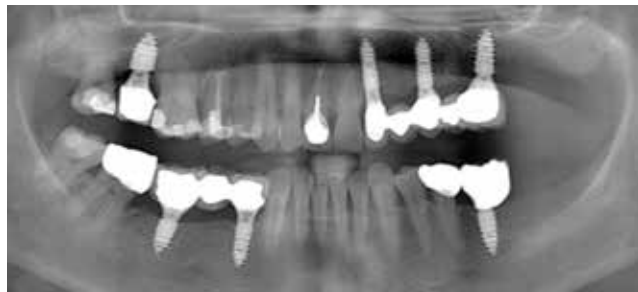
20.05.2014



19.09.2014



20.03.2015

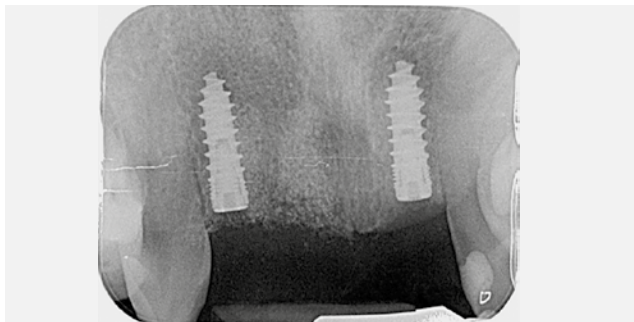


20.11.2020

Case 11: immediate implantation  
12 + 21



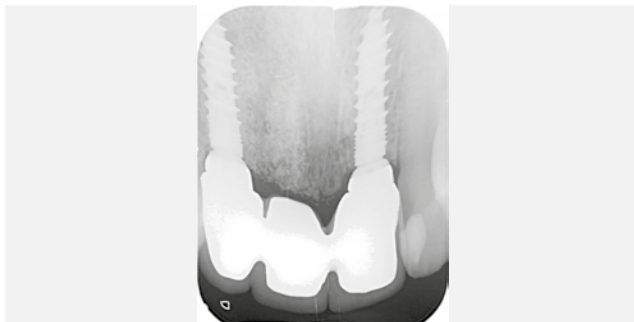
17.02.2015



28.04.2015



28.08.2015

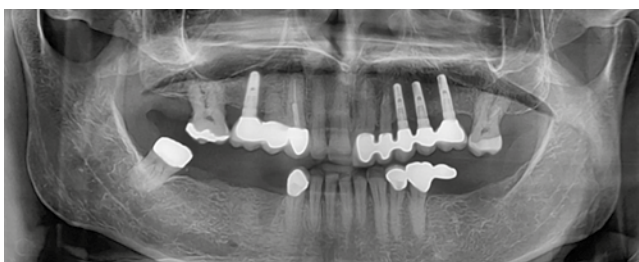


03.01.2018

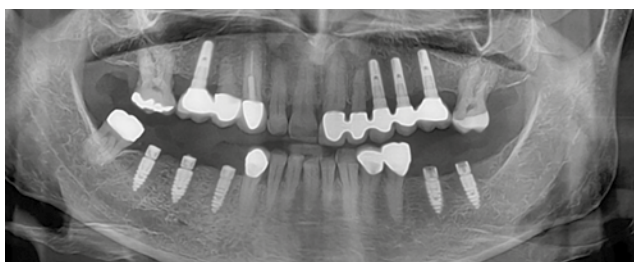


17.04.2020

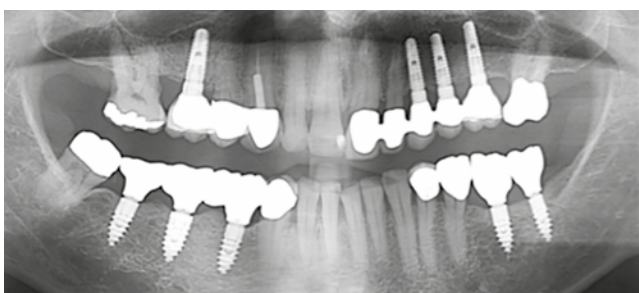
Case 12: Implantations 35, 36, 44-46



18.08.2011



23.09.2011



20.03.2015



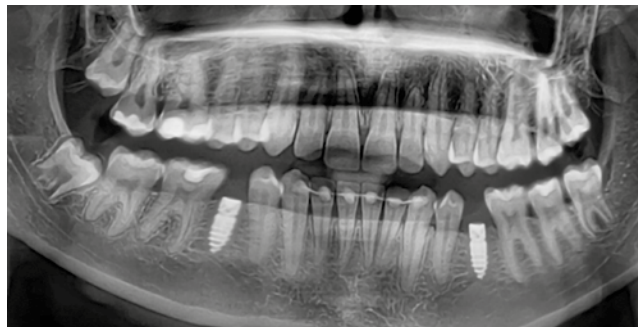
24.04.2020



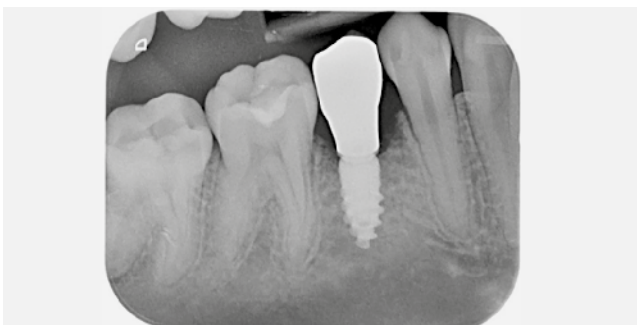
Case 13: Implantation 35 + 45



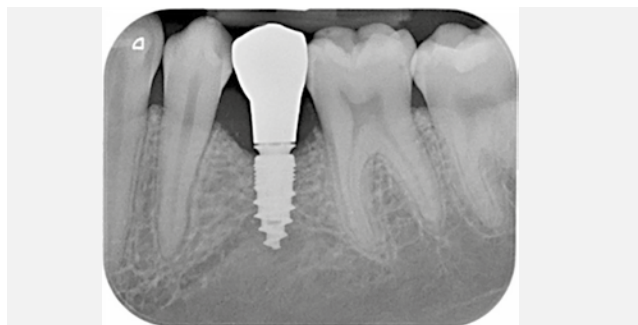
08.11.2011



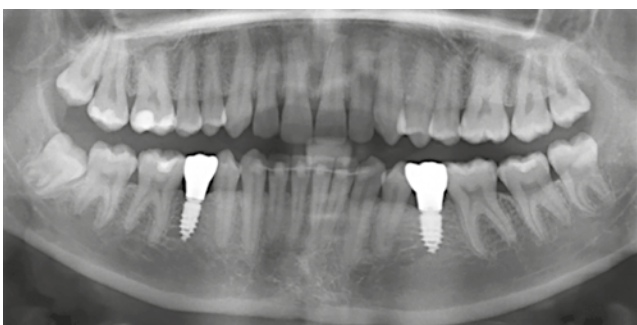
10.02.2012



15.09.2017

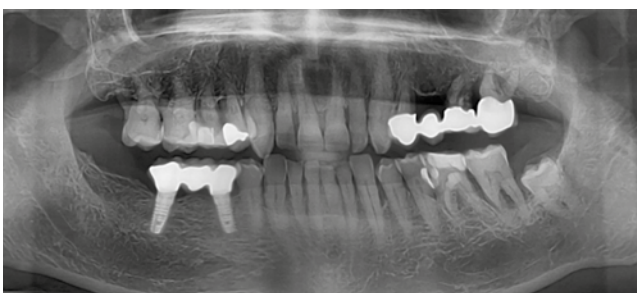


15.09.2017

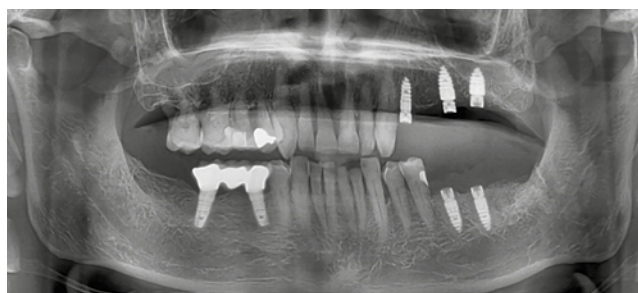


18.06.2021

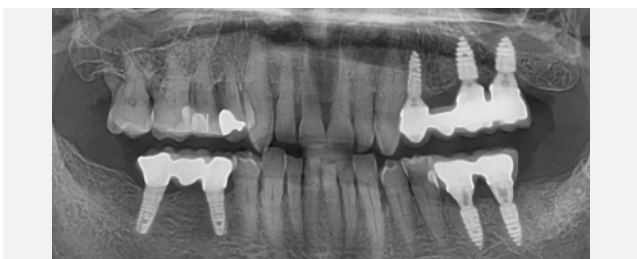
Case 14: delayed immediate implantation 24, 26, 27 + 36, 35



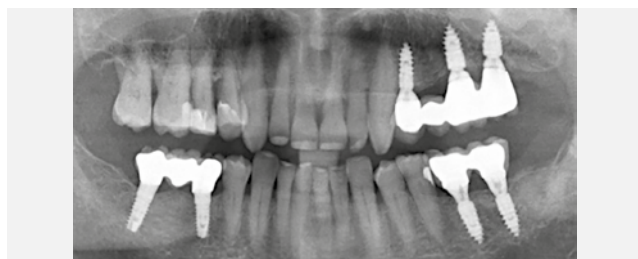
20.05.2016



16.09.2016



22.03.2019

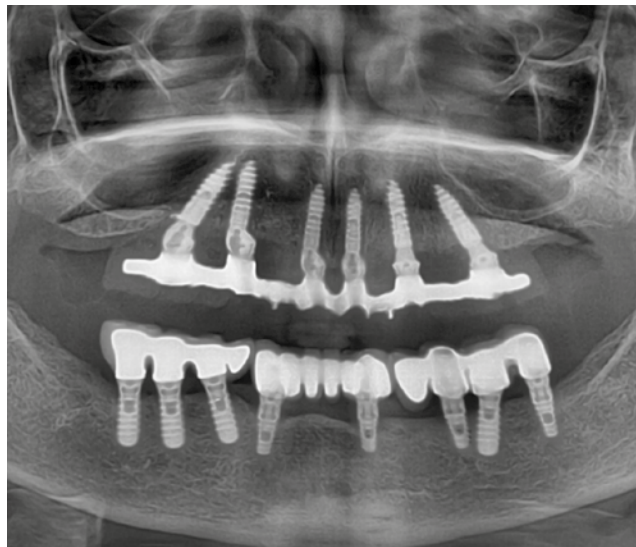


15.09.2021

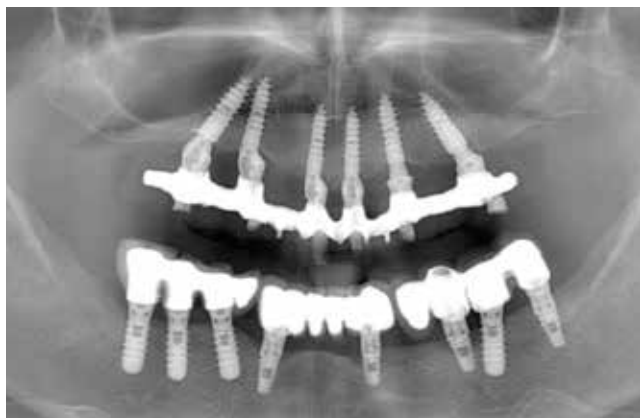
Case 15: implantations 15, 13, 11,  
21, 23, 25 for "All-On-6" concept



13.06.2017

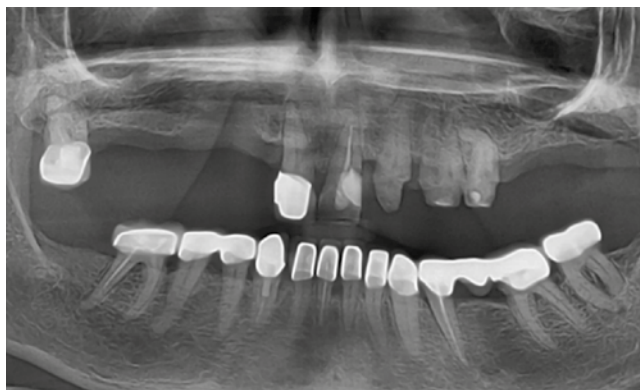


11.02.2018

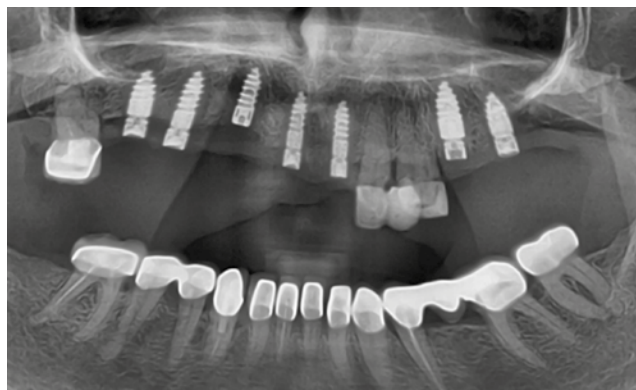


11.03.2021

Case 16: Implantations 16, 15, 13,  
11, 21, 25, 26



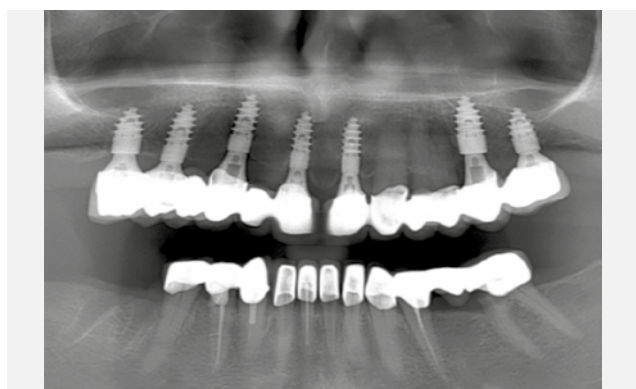
18.12.2014



13.02.2015

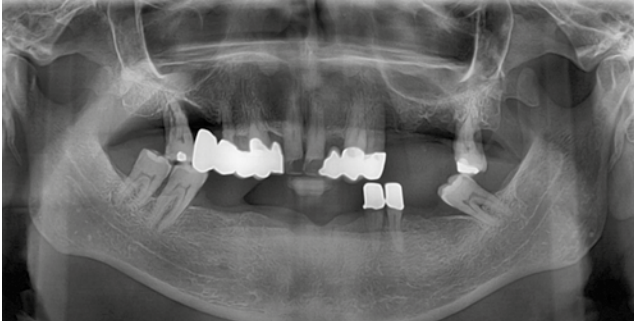


13.11.2018

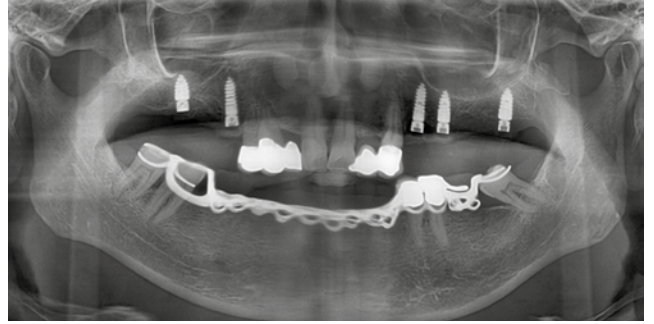


10.03.2021

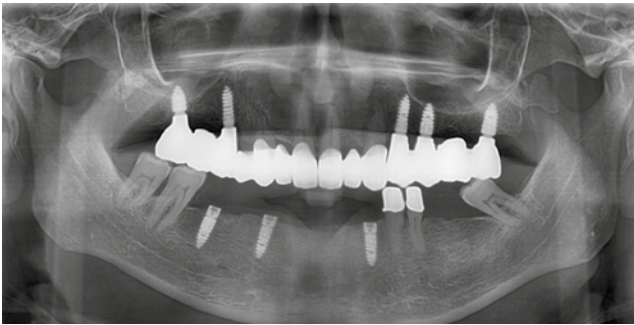
Case 17: implantations 17, 15, 24,  
25, 27, 32, 43, 45



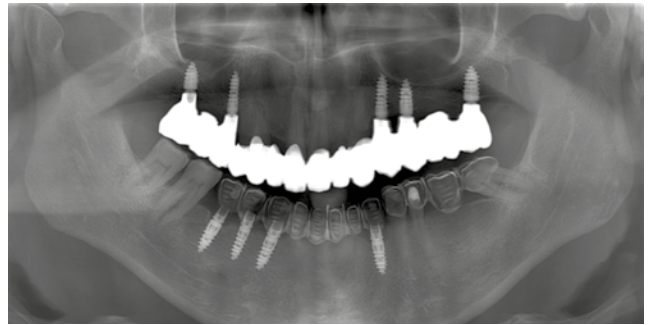
08.09.2011



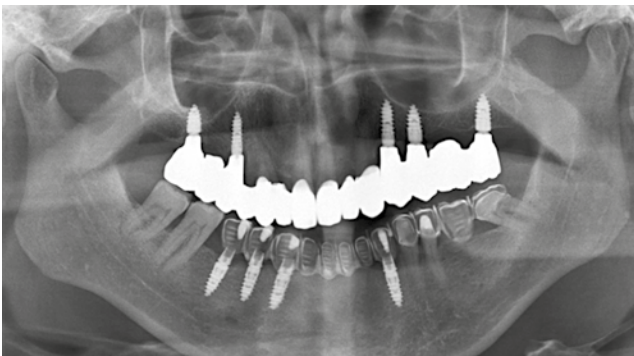
14.10.2011



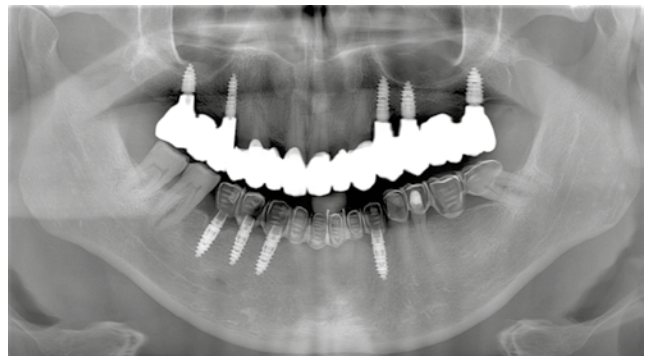
25.01.2012



01.06.2015



27.07.2018

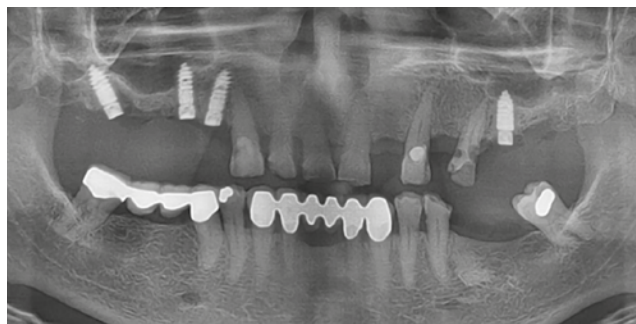


19.03.2021

Case 18: implantations 17, 15, 14, 26  
+ 45, 46, 47



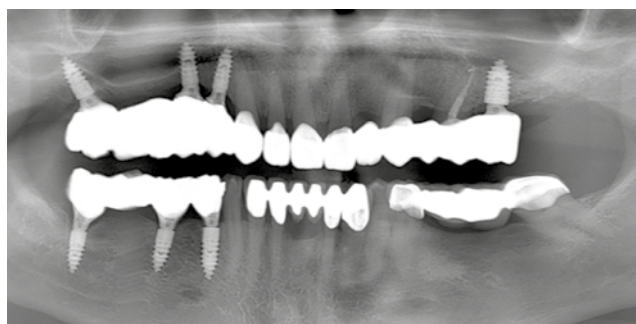
14.02.2013



03.05.2013

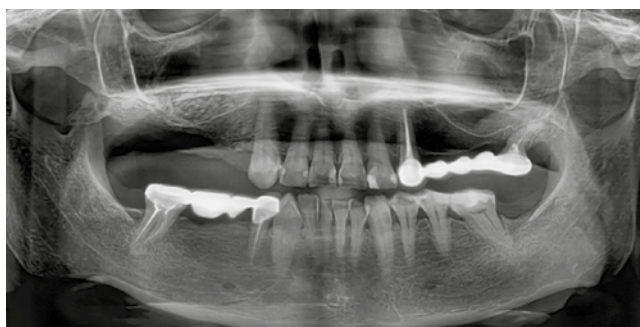


22.08.2018

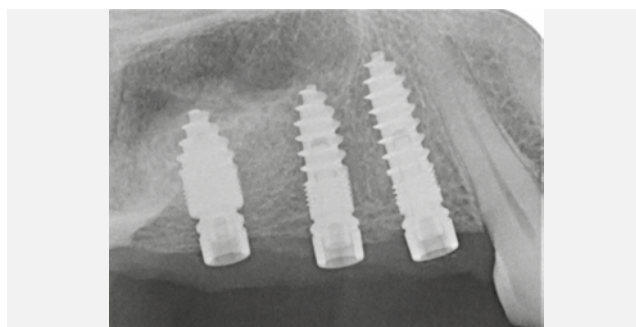


18.08.2021

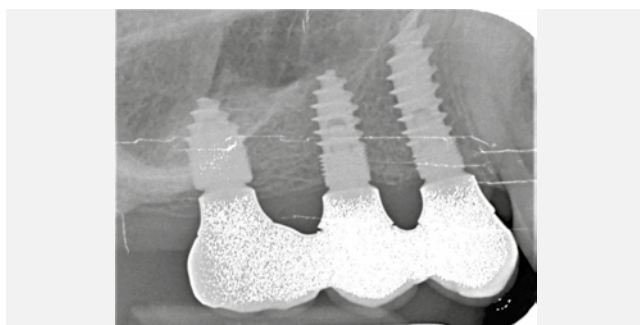
Case 19: implantations 16, 15, 14 +  
immediate implantations 23, 24, 25,  
27



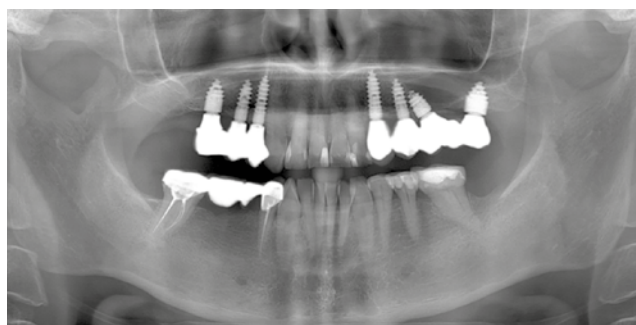
02.02.2015



13.05.2015

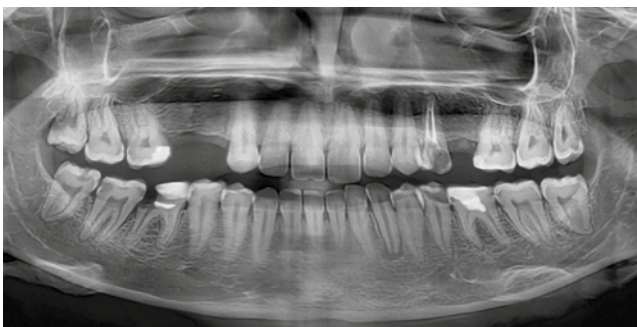


20.09.2017



20.10.2020

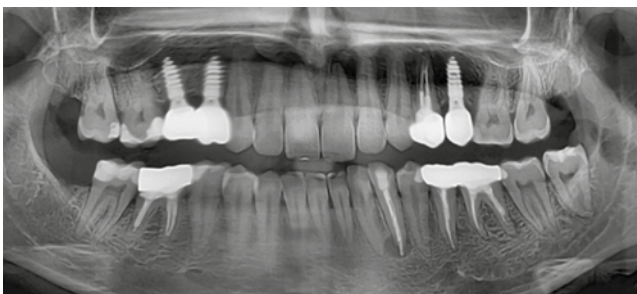
Case 20: implantations 15, 14, 25



01.07.2014



13.06.2014



12.12.2016

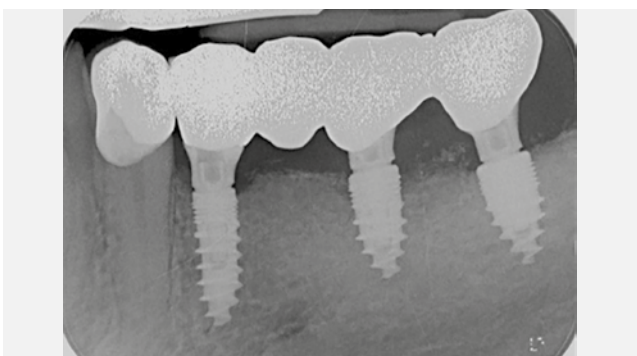
Case 21: Implantations 34, 36, 37



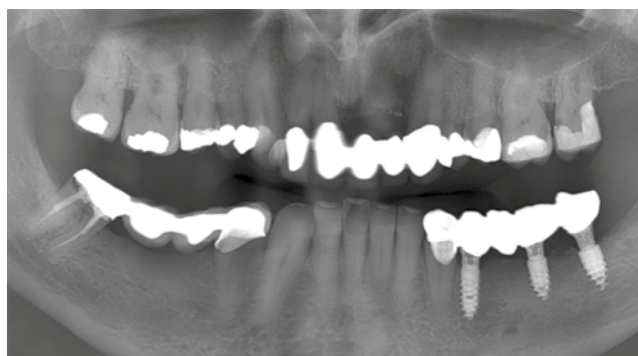
01.03.2012



18.05.2012



07.11.2017

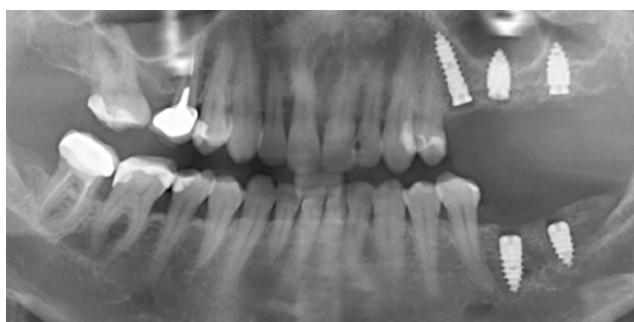


08.06.2021

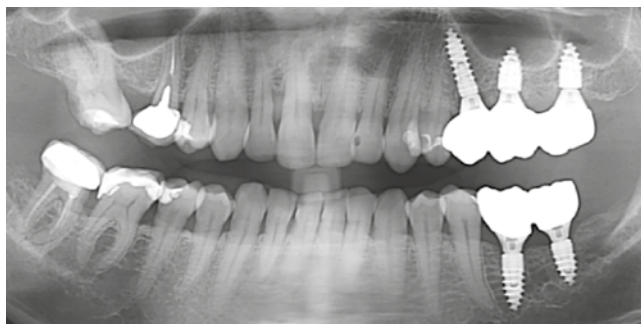
Case 22: implantations 25, 26, 27,  
37, 36



17.12.2013



14.02.2014



15.09.2021



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