A study of the bone healing kinetics of plateau versus screw root design titanium dental implants

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Abstract

Objective: This study was designed to compare the bone healing process around plateau root from (PRF) and screw root from (SRF) titanium dental implants over the immediate 12 week healing period post implant placement.

Material and methods: 32PRF and 32SRF implants were placed in 8 beagle dogs at 12, 8, 5 and 3 weeks prior to enthanisation using a bilaterally balanced distribution. Undecalcified ground sections were prepared from the biopsies taken and histometric measurements of bone implant contact (BIC) and bone area fraction occupancy (BAFO) were made on the middle 5 mm portion of each 8 mm implant root length.

Results: The analysis showed that although measurements of bone to implant contact (BIC) and bone area fraction occupancy (BAFO) tended to be greater for the SRF implants at all four time points, the differences in measurements between implant types did not reach statistical significance (P = 0.07, P = 0.06). The effect of time on BIC and BAFO was found to be strongly significant for both implant types thus indicating a statistically significant increase in BIC and BAFO overall with time (P = 0.004, P = 0.002). Furthermore, both PRF and SRF implants behaved similarly over time with measurements of BIC and BAFO progressing in parallel. Histomorphologic analysis of these sections demonstrated the prominent role of woven bone (callus) in the bone healing process around PRF implants.

Conclusion: The results can be interpreted to indicate a comparable development of secondary stability for both PRF and SRF implant designs. However, as these parameters reflect the structural connection between implant and bone and not the functional properties of the bone to implant interface, they cannot be regarded as comprehensive measures of osseointegration. This particularly relevant given the reduced load bearing capacity of woven bone.

In the intervening years since Bränemark et al.’s (1969) seminal publication on the osseointegration of screw root form [SRF] titanium dental implants, these implants have become widely accepted. The bone healing process around SRF endosseous dental implants has been well documented in scientific literature [Adell et al. 1981; Albrektsson et al. 1981; Roberts 1988]. A wide variety of different endosseous implant morphologies have been tried over the years but many have fallen out of use due to an inability to satisfy established success criteria [Albrektsson et al. 1986; Smith & Zarb 1989; Albrektsson & Sennerby 1991; Fiorellini et al. 1998].

The plateau root form [PRF] endosseous dental implant, also referred to as a finned or serrated implant, has been employed in a number of guises since the early 1970s [Driskell et al. 1971; Chess 1990]. It is currently available as Bicon + dental im-
plant. The PRF dental implant differs from SRF dental implants in having a series of separate circumferential fins spaced along the bone-interfacing portion of the implant. One potential significance of this design is that there is a large space available between the fins of the implant in which a blood clot can form and in which woven (callus-like) bone can develop. It has been suggested that the vascularisation and filling sequence in this space could result in a relatively rapid biomechanical stabilisation of the implant (Lemons 2004).

The premise that the bone healing process around PRF implants differs from that around SRF implants is based on a very small body of literature (Lemons 2000; Coelho & Suzuki 2005). Berglundh et al. (2003) and Buser et al. (2004) have provided histological evidence of prominent woven bone formation and maturation within experimental wound chambers cut into the surface of non-PRF implants. The aim of this study was to perform a histological comparison of the bone healing process around PRF and SRF implants over the initial 12-week healing period post implant placement using an in vivo beagle model.

Materials and methods

The devices

The SRF dental implants used in this study were Standard Plus Narrow Neck design implants (Straumann® AG, Basel, Switzerland). The implants were manufactured from commercially pure titanium (CPTi) with sand-blasted and acid-etched root surface (3.3 mm in diameter × 8 mm in length) and a machined collar (1.8 mm) and external octagon attachment [Fig. 1]. The vertical distance between root threads was 1 mm and the maximum distance between the central root shaft and tip of the root threads was 0.25 mm. The machined collar flared out to a diameter of 3.5 mm at its upper border. A machined narrow neck cover screw was utilised post implant placement.

The PRF dental implants used in this study were custom manufactured by Bi-con® Inc. (Boston, MA, USA). The implants were manufactured from titanium alloy with a grit-blasted and acid-etched root surface (3.3 mm in maximum diameter × 8 mm in length) and a machined collar (1.8 mm in height × 3.5 mm upper border diameter). The vertical distance between root plateaus was 0.75 mm and the maximum distance between the central root shaft and tip of the root plateaus was 0.75 mm [Fig. 2]. The PRF implants also had an external octagon attachment identical to that present on the SRF implant thus facilitating the use of the same cover screw (Straumann® narrow neck cover screw).

Similar root surface topographies (sand-blasted and acid-etched) and supracrestal morphologies (external octagon and a machined collar 1.8 mm in height × 3.5 mm upper border diameter) were deliberately employed for both implant designs in order to minimise non-root morphology variables.

Experimental animals

The mandibular premolar regions of eight young adult male Beagle dogs were used for the study. The protocol followed was in accordance with the animal research policies of the Department of Health in Ireland.

All surgical procedures were performed under general anaesthesia. Food was withheld for 12 h before each procedure. The animals were premedicated 30 min preoperatively with 0.1 mg/kg of acetyl promazine [Acepromazine maleate BP; Novartis® Basel, Switzerland; 2 mg/ml] by subcutaneous injection. Anaesthetic induction was achieved with 0.05 mg/kg of thiopentone sodium (Rhône Mérieux, Dublin, Ireland) intravenously. An endotracheal tube was inserted and anaesthesia was maintained...
with halothane [Halothane®, Mumbai, India] and oxygen. Local infiltration with 2% Lignospan special used for analgesia and to aid in haemostasis (Lignospan special® Septodont; Mazamet Cedex, France).

The mandibular premolars (P₁, P₂, P₃, P₄) were removed on both sides. To facilitate the removal, a buccal mucoperiosteal flap was raised. The surgical sites were sutured with absorbable sutures [3/0 Vicryl Rapide; Ethicon®, Johnson and Johnson® Langhome, Pennsylvania, USA] and the dogs were allowed to recover from the anaesthetic. They were reinstalled in kennels where they were kept for a healing period of 3 months.

The animals were given 1 ml of carprofen (Rimadyl; Pfizer, New York City, New York, USA; 50 mg/ml) subcutaneously for analgesia at the time of implant surgery and once daily for the next 5 days. In addition, 3 ml of the antibiotic enrofloxacin (Baytril; Bayer AG, Leverkusen, Germany; 50 mg/ml) was administered subcutaneously at the time of implant surgery followed by 150 mg of oral clindamycin (Antirobe; Pharmacia; 150 mg capsules) once daily for the next 5 days.

The animals were reinstalled in kennels after each procedure and were sacrificed by intravenous injection of 200 mg/kg of pentobarbitone [Rhone Merieux] after 24 weeks (12 weeks post initial implant placement session). Block biopsies were harvested and placed in 10% formalin. The specimens were then rinsed in running tap water, trimmed and dehydrated using a graded series of increasing ethanol concentrations. They were embedded in methylmethacrylate without being decalcified.

Tissue blocks were cut in the buccolingual plane parallel to the long axis of the implant into 100 μm thick vertical sections using a slow speed diamond precision saw (Isomet® 1000; Buehler GmbH, Düsseldorf, Germany). The sections were ground and polished to a final thickness of 80–100 μm using a series of silicon carbide coated paper discs on a variable speed polishing wheel [Metprep 10 DVT; Metprep Ltd, Coventry, UK] and surface stained with a 1% aqueous solution of toluidine blue [Donath & Breuner 1982; Rohrer & Schubert 1992].

The slides were analysed using light microscopy 12.5 × (Olympus IX-51 inverted research microscope; Olympus, Tokyo, Japan). Digital micrographs of each specimen were captured [Olympus DP-70 digital camera, Olympus]. The micrographs (*TIF format), were transferred to a personal computer and histomorphometric analysis was performed using analysis software (AnalySIS®; Soft Imaging System GmbH, Münster, Germany).

Histomorphometric analyses were carried out for each implant with the measurement of bone-to-implant contact (BIC) and bone area fraction occupancy (BAFO). A rectangular grid 5 mm (5000 μm) in length was superimposed on the middle 3 mm portion of each 8 mm implant body (Figs 5 and 6). Both BIC and BAFO were measured within the confines of this grid, thus

### Table 1. Implant location per animal, time-point, site and implant morphology

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PRF, plateau root form; SRF, screw root form.

*Weeks pre-euthanisation.

Buccal and lingual soft tissue flaps were raised at the experimental sites. A minimum of 4 mm of vital bone was maintained between implant surfaces and the flap surgery was carried out as conservatively as possible in order not to affect the healing around the previously placed implants. The SRF implant osteotomies were prepared according to the Straumann® surgical protocol. The sterile SRF implant was placed manually to engage the full depth of the 9 mm osteotomy. This resulted in the subcrestal positioning of the sand-blasted and acid-etched root surface (8 mm) and a portion (1 mm) of the machined collar. A cover screw was then placed followed by careful flap repositioning using 3.0 absorbable sutures [Vicryl Rapide, Ethicon®, Johnson and Johnson® Langhome, Pennsylvania, USA].

The PRF implants were placed according to the Bicon® surgical protocol with the exception of depth of placement. Conventional Bicon® implants (have a sloping shoulder, internal taper connection and no machined portion of each 8 mm implant body (Figs 4a–f). They were given 1 ml of carprofen (Rimadyl; Pfizer; 50 mg/ml) subcutaneously for analgesia at the time of implant surgery and once daily for the next 5 days. In addition, 3 ml of the antibiotic enrofloxacin (Baytril; Bayer AG, Leverkusen, Germany; 50 mg/ml) was administered subcutaneously at the time of implant surgery followed by 150 mg of oral clindamycin (Antirobe; Pharmacia; 150 mg capsules) once daily for the next 5 days.

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*Weeks pre-euthanisation.
excluding measurements in the apical and coronal 1.5 mm of each implant body. The BIC values generated represent the total BIC (mm) within the grid expressed as a percentage of the total implant perimeter (mm) within the grid (Fig. 7). The BAFO values generated represent the area of bone (mm$^2$) within the retentive features of each implant expressed as a percentage of the total area (mm$^2$) between the retentive features of each implant (restricted to the middle three interthread/interplateau areas in each grid) (Fig. 8).

Data analysis
The data was analysed using the statistical software package JMP 5.2 (SAS Institute, Cary, NC, USA). Repeated measures analysis of variance tests were employed (MANOVA) to investigate (1) the effect of implant type on BIC and BAFO over time, (2) the effect of time on BIC and BAFO and (3) the effect of the interaction of implant type and time on BIC and BAFO.

Results
For the calculation of %BIC and %BAFO, a total of 48 of the 64 implants originally placed were included in the results ($22 \times $ PRF, $26 \times $ SRF). Sixteen implants were excluded from the results ($10 \times $ PRF, $6 \times $ SRF). From the excluded implants 12 ($10 \times $ PRF, $6 \times $ SRF) that were placed in the Pt ridge region were in contact with the root of the adjacent cuspid tooth. This resulted in the fibrous encapsulation of the implants and a failure of osseointegration. The remaining four ($4 \times $ PRF) all failed probably due to insufficient primary stability. The failed implants were distributed evenly between the different healing times (five at 12 weeks, four at 8 weeks, three at 5 weeks and four at 3 weeks).

The results for percentage BIC and BAFO for PRF and SRF implants in groups corresponding to the four time-points of implant insertion are presented in Figs 9a–e and 10a–e, respectively. The BIC results recorded for the SRF implants were higher at four time-points, increasing from 70.9% at 3 weeks to 89.6% at 12 weeks. For the PRF implants, they increased from a low of 57.5% at 3 weeks to a high of 84.4% at 12 weeks. The BAFO results recorded for the SRF implants were higher at all four time-points, increasing from 58.2% at 3 weeks to 74.7% at 12 weeks. The results for the PRF implants increased from a low of 43.1% at 3 weeks to high of 63.7% at 12 weeks.

Regarding the effect of time on BIC and BAFO, it was shown to be statistically significant (d.f. = 3, F-ratio = 33.2426, $P = 0.004$) and (d.f. = 3, F-ratio = 17.1611, $P = 0.0024$), respectively. This indicates a statistically significant increase in BIC and BAFO overall (for both implant types) with time.

The effect of implant type on BIC and BAFO approached but fell just short of being significant (d.f. = 1; F-ratio = 4.2865, $P = 0.0722$) and (d.f. = 1; F-ratio = 4.5230, $P = 0.0661$), respectively. Non-significant was also the effect of the interaction of implant type and time, which indicates that with regard to BIC and BAFO, both types of implant behaved similarly over-time.

Discussion
The bone healing process around SRF implants has been well documented and a wealth of supporting histological evidence exists as to the role of bone modelling and remodelling at the bone to implant interface (Brånemark et al. 1969; Albrektsson et al. 1981; Roberts 1988; Suzuki et al. 1997; Davies 2005). However, the premise
that the bone healing process around PRF implants differs from that around SRF implants is based on a very small body of literature.

Lemons (2002) performed a qualitative histological analysis of PRF and SRF dental implants explanted from humans for reasons other than a failure of osseointegration. Based on his findings, Lemons (2002) suggested that the bone healing process around PRF implants differs from that around SRF implants, in having a greater contribution from woven bone formation. However, as all the implants had been in clinical function for more than a year, Lemons (2002) had no histological evidence of woven bone infill in the early bone healing phase (0–3 months). Unlike his work, this study provided definitive histological evidence of the significant role of intramembranous ossification in the osseointegration of PRF implants in the 0–3 month healing phase. Furthermore, through qualitative histological analysis, it was possible to show that intramembranous ossification played a more prominent role in the peri-implant bone healing process around the PRF implants than it did for the SRF implants that were placed at the same time-points.

This study was unique in having the expressed aim of comparing the bone healing process in PRF and SRF dental implants. The parameters BIC and BAFO are well-established measures of osseointegration in scientific literature.
It is well recognised that the proportion of BIC depends on a number of factors including implant material, surface characteristics, surgical technique and site, time and implant design (Albrektsson et al. 1993). The study was structured so as to minimise the effect of all but the last of these variables, as the effect of implant design was the focus of our attention. While the PRF and SRF implants were made from titanium alloy and CPTi, respectively, they both had a sand-blasted and acid-etched surface. The authors are unaware of studies that compare similar implant types within the study that is of concern. The clinical validity of such a comparison may be duly questioned on the grounds that the morphology of SRF and PRF implants is so different. Given the considerably greater surface area and inter-plateau area of a PRF implant to a similar length SRF implant, it may be questioned whether an equivalent %BIC or %BAFO is clinically necessary. In order to answer these questions, it is worth considering the definition of osseointegration as ‘a direct structural and functional connection between ordered living bone and the surface of a load carrying implant’ (Brånemark 1983). While BIC and BAFO are established histologic measures of the structural connection between implant and bone and interthread bone density, they do not provide a direct measure of the functionality of that connection. In isolation, they are not comprehensive measures of osseointegration. The quantification of the functionality of the connection between implant and bone is a more complicated and situation-dependent calculation, requiring other scientific tests such as mechanical testing and finite elements analysis. Furthermore, as woven bone has inferior load-bearing properties to ordered lamellar bone (Roberts et al. 1987), a calculation of the proportion of woven bone to lamellar bone in the interthread/plateau area might be appropriate as a further histomorphometric measure in this study.

This study showed that, even though higher values were recorded for measurements of BIC and BAFO for the SRF implants at all four time-points, the difference between these measurements for SRF and PRF implants was not significant. Given that PRF implants initially demonstrate a stronger performance on the part of the PRF implants was not significant. Given that PRF implants initially demonstrate a less intimate relationship with the osteotomy than SRF implants, this represents the strong performance on the part of the PRF implants in achieving effective parity for these parameters. As there was no statistically significant difference in the BIC and BAFO trends for both implant types
between weeks 3 and 12 [the measurements increased in parallel], it can be speculated that there was more rapid bone formation around the PRF implants between insertion and the 3-week time-point. In the absence of BIC and BAFO measurements immediately after insertion, this cannot be scientifically verified. This may be considered a limitation of this study.

The results of this histomorphometric study of the bone healing process around PRF and SRF dental implants will contribute to the very limited body of scientific evidence available to date on this topic. The histomorphometric calculations, BIC and BAFO, provide a record of the structural connection between implant and bone at a light microscopic level. While this may be accepted as evidence of the rapid stabilisation of PRF implants, it must be recognised that, by definition, osseointegration requires both a structural and a functional connection between implant and ordered living bone (Brånemark 1985). In an age where the traditional 12-week healing period between the placement and functional loading of SRF implants is steadily decreasing, the poor load-bearing capacity of immature woven bone must raise questions as to the relative performance of PRF implants under such conditions. Further research will be necessary to investigate the relative functionality of the bone to implant connection for PRF and SRF implants in the traditional 12-week healing period post implant placement. This might involve a similar study design with mechanical testing of unloaded PRF and SRF dental implants.

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References


