

A Six-Year Prospective Study of 620 Stress Diversion Surface (SDS) Dental Implants

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ABSTRACT

A 6-year prospective study was conducted to assess the clinical success rates and crestal bone response of a dental implant system with a stress diversion design. Mathematical modeling, digital radiography with applied isodensity and Finite Element Analysis were used to highlight the effect of the stress distribution design. A total of 386 hydroxyapatite coated prototype, and 234 commercial grit-blasted external hex implants were placed in virgin bone, as well as various grafted maxillary regions, with 36% of the posterior implants being immediately loaded. Prototypes achieved 96.6% survival over a 3 year period. The grit blasted implant; placed from 2000 to 2003, showed a 95% survival rate. There were no significant changes in crestal bone levels after the first 12 months of prosthetic loading. Engineering evaluations suggested that undesirable stresses were distributed from the crest of the ridge down through the center of the implant body.

Key Words: clinical trial, dental implants, Resorbable Blast Medium, Finite Element Analysis, Stress Diversion Surface, Isodensity Radiography.

INTRODUCTION

The physiologic response of crestal bone levels surrounding an osseointegrated implant can be separated into three general phases.

The first phase is associated with primary site preparation, and has been well documented in the literature¹. Crestal bone loss during the first phase has been attributed to poor surgical technique, inadequate quality and quantity of bone, or a pre-existing osseous defect not previously controlled^{2,3}.

The second phase would be described as the uncovering phase. Crestal bone loss associated with this surgical phase is scarcely reported in the literature, due to the noninvasive nature of the procedure.

The third phase is that of prosthetic loading and function. Crestal bone loss in this phase is multi-factorial and well reported in the literature^{4,5,6}. These “ailing and failing” implants have introduced a new discipline within the field of implant dentistry, attempting to reverse or repair the obvious clinical problems of crestal bone loss^{7,8}.

Reports of excessive crestal bone loss have been suggested to be a result of plaque-induced peri-implantitis⁹ or occlusal overload, the latter producing excessive stresses in the crestal bone surrounding the implant^{10,11}. Publications demonstrating bone loss at the crestal ridge of implants associated with microbial infections, noted that it occurred only after occlusal overload^{12,13}. One study, utilizing a primate model, demonstrated that occlusal overload was more responsible for crestal bone loss than plaque accumulation, by more than a three-fold difference¹⁴. Both photoelastic and Finite Element Analysis (FEA) demonstrate that the highest concentration of stress at the implant/bone interface

occurs within the area of crestal bone^{15,16,17}. High crestal implant-to-bone stresses may also accelerate the accumulation of plaque in a similar way that occlusal stresses increase plaque accumulation around natural dentition¹⁸. The increased threshold to tactile perception, associated with up to fifty times greater force in dental implants, subjects the dental implant to the increased risk of occlusal overload^{19,20}. Desmodontal structures refer to all physical, chemical and physiological structures associated with the myriad of events within the dental and periodontal attachment apparatus. This system is a complex sympathetic and parasympathetic coordination of all neuro-muscular structures of the myofacial system, designed to prevent overload and harmful assaults to the natural dentition²¹. This is missing in the peri-implant region due to the missing desmodontal structures^{22,23}. Their absence in dental implants enhances harmful stresses, and puts dental implants at greater risk for crestal bone loss than natural teeth^{24,25}. Late failure of osseointegrated dental implants is most commonly the result of bone loss at the crest of the alveolar ridge²⁶. The clinical reality of this ever-present problem in implant dentistry has led to a myriad of solutions.

In recent years, studies have focused on improving the clinical success of osseointegrated implants by evaluating the effect of design and shape.^{27,28,29} Bioengineering has played a major role, with its increasing effectiveness in determining the optimum design for implant configuration and the prosthetic superstructure^{30,31,32}. Geometric analysis of the dental implant has shown a threaded architecture to be a more favorable design for ideal stress distribution, when compared to non-threaded implants³³. Mathematical and mechanical modeling of the bone surrounding an implant can provide theoretical information concerning the effects of implant geometry and the bone environment.

There is an ideal envelope of stress that encourages a physiologic osteoblast response in bone remodeling, and a corresponding relative magnitude of stress associated with the process of bone resorption and maturation³⁴. However, it has yet to be scientifically and clinically proven which thread configuration would be most favorable in evenly distributing the stress of occlusion. The objective of this prospective study is to clinically analyze the effectiveness of a dental implant with a stress-diversion design in minimizing crestal bone loss, and enhancing bone maturation below the bone crest.

METHOD AND MATERIALS:

The SDS implant (Stress Diversion Surface- ACE Surgical, Brockton, Mass.), shown in Figure 1, is made of type IV commercially pure titanium. This surface is treated with a resorbable blast medium (RBM)³⁵. All prosthetic connections are the industry standard external hex (Figure 1A). The transition from the prosthetic connection to the thread configuration is characterized by a highly polished 0.7mm titanium surface (Figure 1B). The coronal, or proximal 6.5mm of the thread (Figure 1C), is the stress diversion thread configuration. It consists of one continuous thread that progressively changes the thread angle and inner minor diameter, while holding the major diameter constant. This thread is mathematically designed to divert the stress of occlusal load away from the crest of the ridge, and into the implant body. The thread transitions into a continuous self-tapping tapered design.

Tables 1 and 2 demonstrate the distribution of implant placement in bone after surgical placement. At least 70% were placed in the first two years of the study. The majority of the implants were 15 mm in length (64.1%), with the 13 and 10mm lengths representing 19.6% and 16.3%, respectively, of all implants placed. The largest numbers of implants

(138, or 59%) were loaded immediately, while 96 (41%) were unloaded. There was some variability in the depth to which the head of the implant was placed relative to the crest of the ridge. In 158 implants (67.5%), the top of the implant neck was crestal, while 76 (32.4%) were up to 1-1.5 mm above the crest.

A minority of the implants (91, or 38.8%) was placed into native bone with no additional grafting, while the remainder were placed with the addition of some type of augmentation material; either 1) an amino acid peptide and anorganic bone mineral (PepGen P-15, Dentsply Ceramed Dental, Lakewood, CO.) alone, 2) a combination of this anorganic bone mineral and deproteinized mineralized cancellous bone allograft (Puros, Zimmer Dental, Carlsbad, CA.), 3) an allograft mixture with autologous Platelet Rich Plasma (PRP), or 4) subantral augmentations with a combination of PRP with a freeze-dried demineralized bone allograft, deproteinized human cortical bone (banked tissue) and a microfibrillar bovine collagen hemostatic agent (Collastat OBP™, Integra Corp, Plainsboro, NJ) (Table 2). Implants were placed in all jaw locations with the anterior and posterior mandible, as well as the anterior maxilla, each containing close to 50 implants. A larger number, or 92 (39%), were placed in the posterior maxilla. Of these 92 implants, 51 (22%) were placed in a subantral augmentation. It is also noteworthy that 36% of these posterior implants were immediately loaded

Each implant in the study was documented radiographically, and positioned by means of a standard Rinn Kit, with an assigned isodensity value of 4. Digital reverse polarization techniques were used to evaluate the qualitative changes in the surrounding bone. Within a given digital radiograph, using constants (the implant body), the relative changes to the surrounding tissues were compared and analyzed by means of the Vix-

Win computer program^{36,37}. In the present program, the same number of pixels/mm² represents an area of “isodensity”, and given the color yellow³⁸. Every area with the same density will show the same color, based on the degree of mineralization. Figure 3 is an example of this type of analysis. Figure 3A shows the digital radiograph of two study implants in the lower left first molar position. This radiograph was taken 18 months after insertion of the implants and 12 months after occlusal loading. The bitmap format of the digital radiographs allow for standardization of sequential radiographs over the duration of the study. Note also the isodensity mapping of similar areas on the adjacent teeth. Figure 3B is the same radiograph with isodensity mapping of a different region of the bone surrounding the implant. Figure 3 also shows a comparable isodensity mapping for a conventional screw type implant, depicting the crest of the ridge (C) and the remainder of the implant body (D), after two years in function.

Mechanical modeling of implant geometry and bone environment made use of the modeling technique known as Finite Element Analysis (FEA), and was accomplished by using the Ansys 5.3 software program (Ansys Corporate, Canonsburg, PA), a commercially available FEA code.^{39,40}

RESULTS

There were a total of 620 implants placed over a six-year period (Figure 2). From 1997-2000, a prototype design was used. This implant was coated with hydroxyapatite, and was machined without the continuous helical thread configuration, while still maintaining varying angles of incidence. Out of the 386 prototypes placed, 13 (3.4%) have been removed due to total implant failure. Four of these 13 implants were removed

within the first month due to extremely poor quality of bone. Five of the failing implants were removed due to post-surgical infection related to immediate extraction. The remaining implants were removed as a consequence of premature overload associated with immediate placement and habitual grinding. The remaining implants demonstrated consistent bone maintenance and associated circumferential maturation, as opposed to crestal bone loss. In the surviving 96.6% of the implants, there were no signs of crestal wear.

The 234 implants placed from the years 2000 – 2003 (Figure 2), were the current commercial design, with an RBM surface treatment. Twelve of these implants were removed. Eight of the twelve failed due to prosthetic overload at the time of immediate placement. Four of the eight failures were in a subantral augmentation of one patient. The failures in this case can be attributed to patient neglect and heavy bruxism. Two failures were implants placed in immediate extraction sites with pre-existing chronic endodontic infections. The remaining two implants were removed due to overload at the time of immediate placement. Three of the 12 failures were in the area of the right lower molars. These implants were prematurely exposed and in very poor quality bone. The failure sites have been grafted with the intent of replacing the removed implants.

Of the 222 implants remaining, 11 (4.4 %) had 1-2 mm of horizontal bone loss after the first 6 months of prosthetic loading. An additional 6 showed similar bone loss after 12 months. There was no change in the radiographic appearance or probing in these implants in the following recall visits. Those implants that showed bone loss were

associated with the group of 91 (Table 2) that were placed in virgin osteotomies. This might suggest that the bone loss was due to surgical trauma, and not prosthetic overload. They also fell into the category of the 76 implants where the head of the implant was placed above the crest of the ridge. Poor initial osseous contouring or quality of bone would most likely explain this bone loss pattern. It is noteworthy that there were no significant changes in the bone level after 12 months of prosthetic loading.

The 3-dimensional FEA of the study implant design, when compared to the industry standard Branemark (Nobel Biocare, U.S.A., Yorba Linda, Ca.) screw implant design, clearly demonstrate that simulated occlusal loads were more evenly distributed in the first 6 to 8mm of the study implant, with diversion of the stress from the crest of the ridge down into the implant body. Figure 4A shows the FEA of a 13 mm study implant (left) and a 13 mm standard implant (right). The surrounding bone matrix was assigned a modulus of elasticity of 8 GPa (Gigapascals) to replicate cortical bone. An *on-axis* compression load of 50 pounds of force (average occlusal load) was introduced with equal parameters in both models. Crestal stress was reduced to 579 pounds per square inch (PSI), depicted by the smaller red band (maximum von Mises stress) around the study implant at the crest of the ridge. The medium stress bands in green on the study implant depict an increase in stress distribution. That is, stress is diverted away from the crest of the ridge and is redirected 2-8 mm below the crest, in even bands, by the implant's thread configuration. The standard implant does not demonstrate this same pattern, as evidenced by the absence of medium (green) stress bands in the center of the implant body, as well as the increased presence of yellow and red stress band at its crestal region. In fact, the increased maximum Von Misses stress band (red) over the 1-2mm

coronal area of the implant has a value of 859 PSI, which is a 48% greater stress application to this area, when compared to the study design. Figure 4B depicts 13 mm study (left) and standard (right) implants. The surrounding bone matrix was again assigned a modulus of elasticity of 8 Gpa. A *transverse* cantilever load of 4.535 inch-pounds, which generates significantly higher application stress, was also introduced with equal parameters to both models. These transverse loads were generated by applying a compressive load of 14.4 pounds, 8mm from the central axis of the implant. Again, Figure 4B depicts the highest red stress band (maximum von Mises stress of 8058 PSI) at the crest of the ridge for the standard implant. The corresponding FEA plot of the study implant, with the same parameters, again shows the minimized crestal red band (maximum von Mises stress of 5626 PSI) and a more even distribution of stress, depicted by the increased blue bands at the implant's thread configuration. Under these transverse loading conditions, the standard implant's maximum crestal stress was found to be 43% higher than the study implant's thread design, which correlates with the on-axis compressive loading results.

DISCUSSION

Knowing the ideal stress requirements necessary for positive bone maturation, and the average range of forces associated with normal implant function, it should be possible to physiologically back-engineer an external implant geometry that diverts the undesirable destructive crestal stress to an area further down the body of the implant. This would create a stress diversion design, which promotes bone growth and maturation under normal loading conditions. Once an implant is integrated, prosthetic loading introduces forces to the implant-bone interface, and the surrounding connective tissue strata, that are

directed through the implant and into the surrounding supportive anatomy. The implant body and its physical properties remain a constant. The supportive anatomy surrounding the dental implant is an ever-changing, visco-elastic medium of multi-potential cells, with the unique ability to change form and structure to accommodate the desired task. This unique collection of connective tissue will be referred to as the peri-implant substrate. The more rigid the connection between the superstructure and this peri-implant substrate, the more force absorbing demand is placed on the implant and its surroundings.

The laws and principles associated with the science of biomechanical engineering govern all aspects of the load dynamics and stress distribution associated with the forces of occlusion and the distribution of those forces. Material science has given us a clear and predictable mathematical explanation of the constants within our prosthetic system. The peri-implant substrate presents with an infinite range of densities, volumes, and modulus of elasticity. This variation also changes with each patient and even with each site in the oral cavity. We are able to establish some ranges in the areas of modulus of elasticity, density and the percent of surface area contact, but these remain variables when mathematically describing the stress distribution around a dental implant. The isodensity mapping of bone surrounding the stress diversion design implants demonstrates a maintenance of a crestal attachment at the very top of the implant after a year in function (Figure 3A), as well as an increased region of bone maturation associated with the stress diversion thread portion of this implant (Figure 3B). This increase in density is an example of the ideal distribution of stress away from the crest of the ridge to a more favorable location 2-8 mm below the crest. The mapping of a standard screw type implant, in Figure 3C, shows the maintenance of a crestal attachment, with similar bone

density at the top of the implant. Figure 3D demonstrates the isodensity mapping of an indiscriminate medullary pattern of bone after two years of occlusal loading. Although there is no sign of crestal saucerization or bone loss, there is also no sign of positive maturation of bone related to functional load as seen in the stress diversion implant design.

Finite Element Analysis (FEA) has demonstrated that the nature of the connection, as well as the properties of the surrounding bone, greatly influence stress distribution³⁷. Several studies have demonstrated that implant geometry and diameter create both positive and negative influences on stress distribution⁴¹. The study dental implant was conceived with the intention of evenly distributing the stresses of occlusion away from the crest of the ridge.

When a prosthesis is put into function, the forces associated with the load dynamics are distributed through the rigid connections and into the surrounding connective tissue. These forces seek out the most rigid material and connection to aid in the support necessary for the particular function. There is material fatigue and component breakage, if given enough time and force to allow this form of damage. In certain cases, bone degeneration often starts at the most rigid connection closest to the origination of the load. Mathematically, it can be demonstrated that the compressive forces introduced by an occlusal load can be up to 5 times greater than the corresponding tensile forces⁴¹. It would stand to reason then, that to reduce the harmful stresses at the crest of the ridge, one would be encouraged to convert these stresses from compressive to

tensile⁴¹. Looking at the mathematical derivation of stress analysis within changing thread values, the ideal angle of incidence and surface area can be extrapolated.

$$\Delta \sum \chi = \Delta \sum \gamma (\frac{1}{2} \text{Sin } \alpha) \quad \text{Equation 1}$$

Where $\Delta \sum \chi$ is the change in the forces introduced in the surrounding bone, and $\Delta \sum \gamma$ is the change in the sum of the forces of occlusion. One can see that as the angle of incidence ($\text{Sin } \alpha$), or the pitch of the thread is increased, the corresponding transverse force into the bone ($\Delta \sum \chi$) is decreased.

Using this formula, the angles necessary to maximize tension at the crest, and minimize compression, can be derived. The distance stress must travel down the body of the implant also counterbalances further stress distribution. This leads to a more even force distribution throughout the first 6-8 mm. This inherently would help compensate for the increased stresses associated with implant dentistry. This stress distribution would also be more likely to yield values within that ideal zone associated with bone development and maturation⁴². The FEA samples indicate a reduction of both maximum stresses, and/or collar region stresses, for various osseous models. This stress distribution indicates that in carrying loads, the study implant uses more of the surrounding bone material than does the standard screw implant. One can see that the combination of high stresses in the collar region, from both compression and bending, can cause resorption and bone loss in this region, leading to "saucerization". Therefore, the study implant, through proper clinical use, could aid in the reduction of these stresses and resulting bone loss. For they're to be a total of only 17 implants with post-prosthetic crestal bone

changes, given the wide range of clinical applications, is of clinical relevance. This would present a total of 91.7 % of the remaining implants, over the given time period, with no sign of crestal degeneration associated with prosthetic loading.

In more evenly distributing the load within the envelope of ideal stress for proper bone maturation³⁴, the study implant design may help to encourage greater bone maturation. Reiger⁴³ has suggested that often an implant's design not only creates maximum stress at the neck and apex, but transfers relatively low stresses to the bone along the body of the implant. This leads to bone loss in the middle of the implant due to atrophy, while at the same time causing bone loss at the extremities, due to excessive stress. In this study, there have been no significant radiographic signs of post-prosthetic crestal saucerization, often seen in other designs within the first 1-2 years of prosthetic loading^{44,45}.

CONCLUSION

Crestal bone loss, and the associated clinical symptoms, is an ever-present problem in the field of implant dentistry. Attempts to prevent or minimize the incidence of crestal bone loss have led to new concepts and designs. The study implant is a change in conventional thread configuration, designed so as to address the factors associated with the forces of occlusion that lead to crestal bone loss. There was a 96% survival rate, with placement in all areas of the jaws, in both grafted and virgin bone. Finite Element Analysis and isodensity radiographic mapping suggest that the stress diversion thread design can lead to more evenly distributed load. While the clinical evaluation of the study implant is still in its early stages, its design concept will shift stresses to more acceptable

parameters, and bone maturation toward higher values than the conventional designs now available.

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TABLE 1							
SDS implants placed per 6 month time period after start of study							
Time (months)							
	0 - 6	6 -12	12- 18	18 -24	24 - 30	30 - 36	Total
No. Implants	58	27	14	82	53	0	234

TABLE 2						
Number of SDS implants in various grafting procedures						
Virgin Bone			Allografts			
No. Implants	No grafting	PepGen ¹	PepGen + Puros ²	PRP ³	PRP + FDDB ⁴ + DHCB ⁵ + Colostat ⁶ (in subantral augmentation)	Total
No. Implants	91	45	7	46	45	234

¹ PepGen P-15™ = Amino acid peptide and anorganic bone mineral

² Puros™ = Mineralized cancellous bone allograft

³ PRP = Platelet Rich Plasma

⁴ FDDB = Freeze dried demineralized human cortical bone (50-100μ)

⁵ DHCB = Deproteinized human cortical bone

⁶ Collastat OBP™ = Microfibrillar bovine collagen hemostatic agent (0.3 gram/5cc)

LIST OF FIGURES

Figure 1. The study implant; A) Standard external hex, B) Polished 0.7mm titanium collar, C) 6.5mm with stress diversion thread, D) Tapered self-tapping implant body, E) “Cutout” in the driving thread.

Figure 2. Number of failures in the six year implant study

Figure 3. Isodensity marking on bone around the, A) study implant configuration at the crest of the ridge and other regions of equal density, B) at the study implant thread portion of the implant, C) the crest of the ridge of a standard screw implant, and D) the remainder of the standard implant body.

Figure 4. FEAs of the study implant and standard screw type implants showing von Mises stresses. In both pairs of schematics the study implant is on the left, and the standard screw implant on the right. Figure 4A depicts on-axis compressive loading of 50 pounds of force (average occlusal load) with equal parameters in both models. Note the reduction in crestal stress, depicted by the smaller red band (maximum von Mises stress) around the study implant at the crest of the ridge versus the standard design. Figure 4B depicts these same implants under off-axis transverse cantilever loading of 4.535 inch-pounds. Again, the study implant shows a decrease in the crestal stress (red) band and the dispersion of these stresses toward the center of the implant.

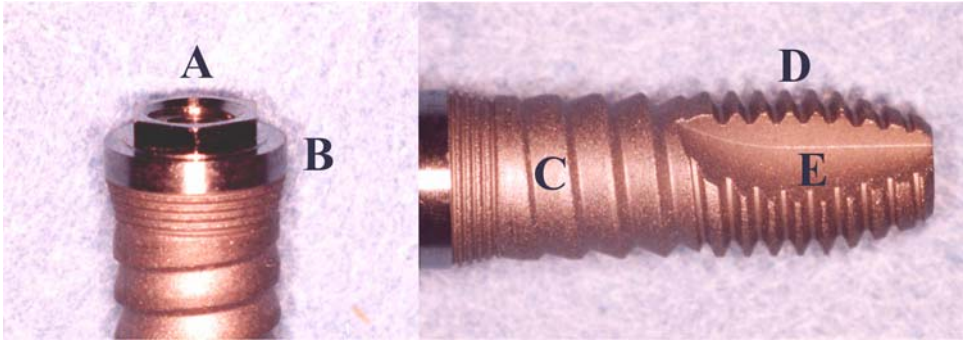
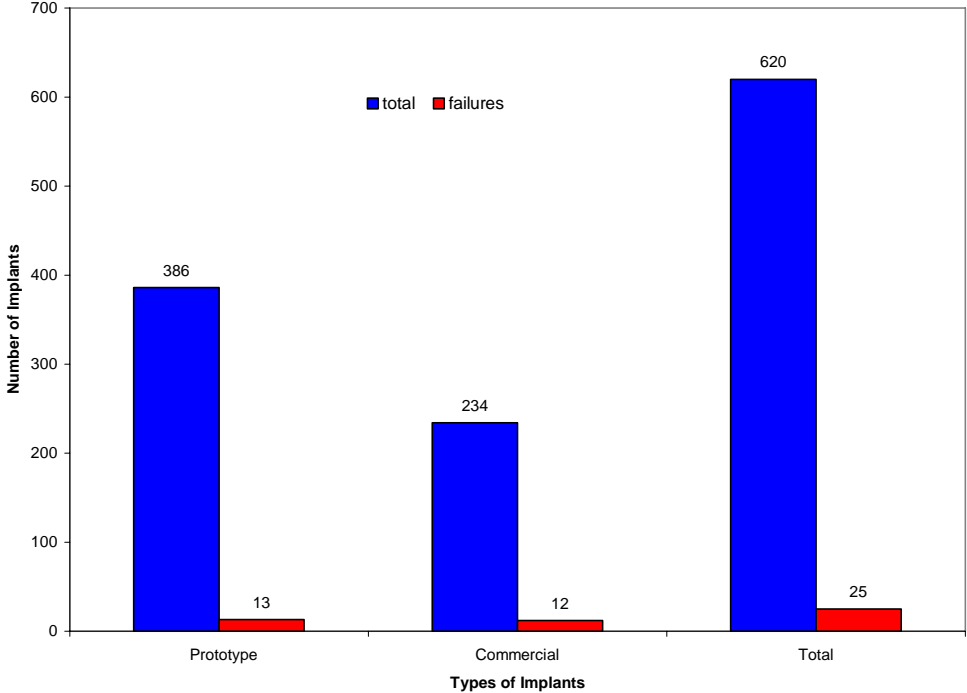


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Figure2. Number of failures in the six-year implant study.



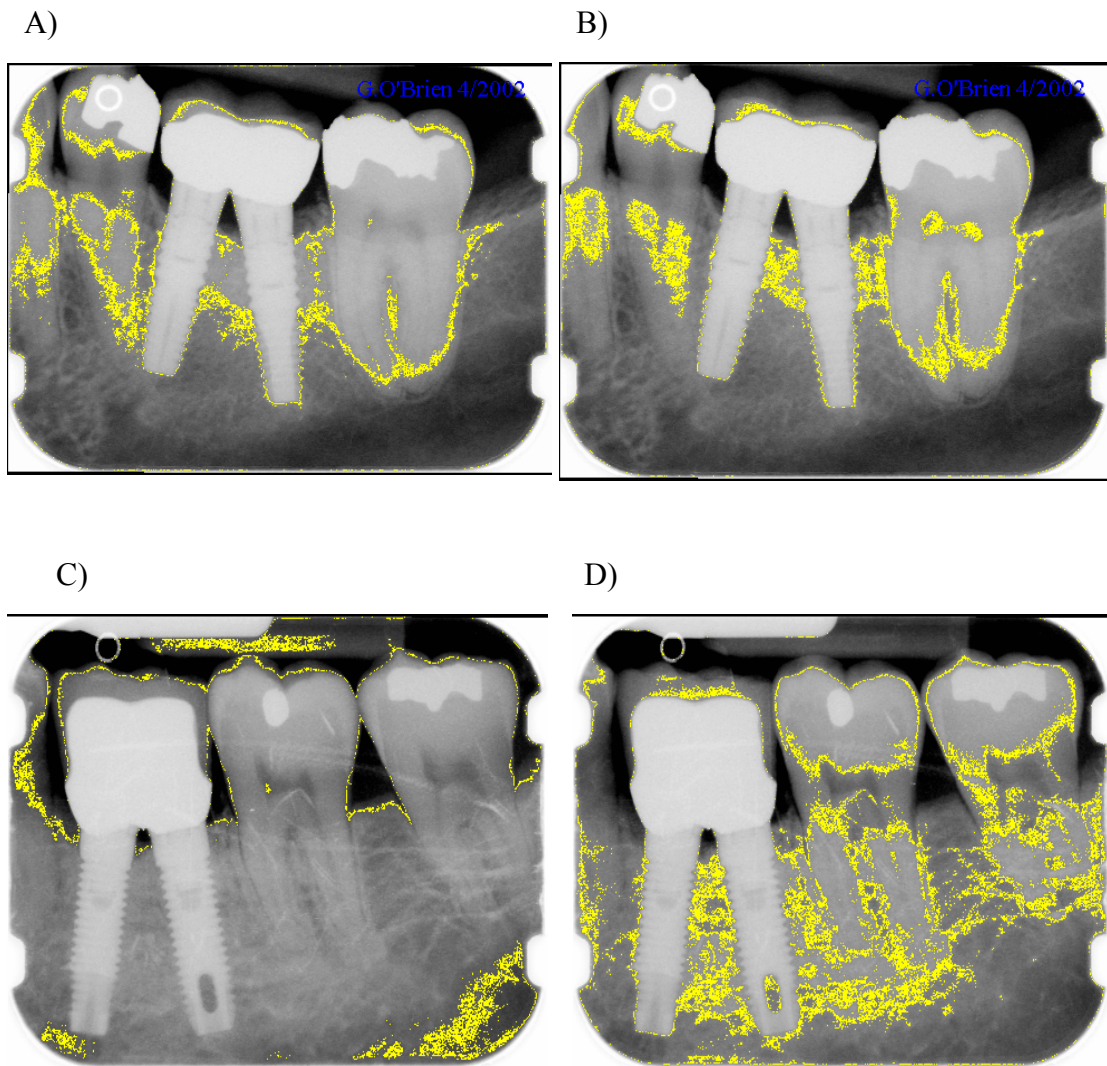


Figure 3. Isodensity marking on bone around the, A) study implant configuration at the crest of the ridge and other regions of equal density, B) at the study implant thread portion of the implant, C) the crest of the ridge of a standard thread implant, and D) the remainder of the standard implant body.

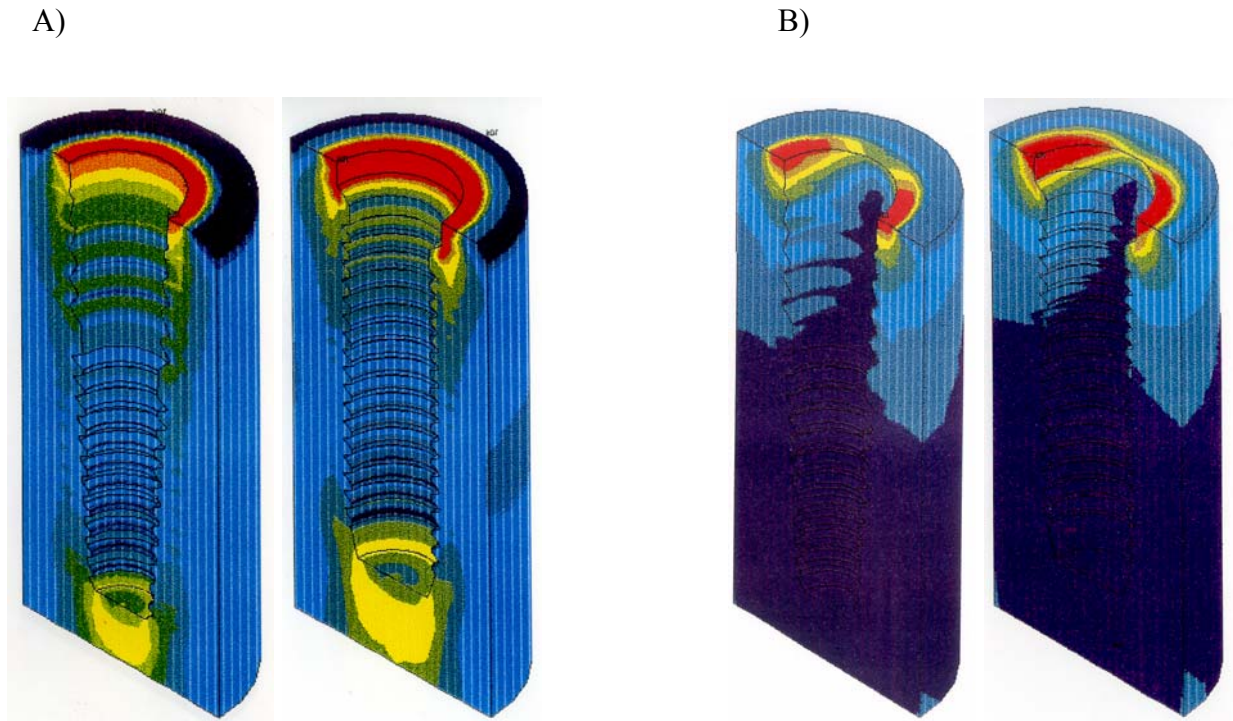


Figure 4. FEAs of the study implant and standard screw type implant, showing von Mises stresses. In both pairs of schematics the study implant is on the left, and the standard screw implant on the right. Figure 4A depicts on-axis compressive loading of 50 pounds of force (average occlusal load) with equal parameters in both models. Note the reduction in crestal stress, depicted by the smaller red band (maximum von Mises stress) around the study implant at the crest of the ridge versus the standard design. Figure 4B depicts these same implants under off-axis transverse cantilever loading of 4.535 inch-pounds. Again, the study implant shows a decrease in the crestal stress (red) band and the dispersion of these stresses toward the center of the implant.