

Flexural strength of mini-implants developed for Herbst appliance skeletal anchorage.

A study in Minipigs br1 cadavers

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Objective: The present study was designed to verify if mini-implant prototypes (MIP) developed for Herbst appliance anchorage are capable of withstanding orthopedic forces, and to determine whether the flexural strength of these MIP varies depending on the site of insertion (maxilla and mandible). **Methods:** Thirteen MIP were inserted in three minipig cadavers (six in the maxilla and seven in the mandible). The specimens were prepared and submitted to mechanical testing. The mean and standard deviation were calculated for each region. A two-way Student's t test was used to compare the strength between the sites. A one-way Student's t test was performed to test the hypothesis. Orthopedic forces above 1.0 kgf were considered. **Results:** The MIP supported flexural strength higher than 1.0 kgf (13.8 ± 2.3 Kg, in the posterior region of the maxilla and 20.5 ± 5.2 Kg in the anterior region of the mandible) with a significantly lower flexural strength in the anterior region of the mandible ($P < 0.05$). **Conclusion:** The MIP are capable of withstanding orthopedic forces, and are more resistant in the anterior region of the mandible than in the posterior region of the maxilla in Minipigs br1 cadavers.

Keywords: Functional appliances. Dental implants. Orthodontic appliances. Orthodontic anchorage procedures. Angle Class II malocclusion.

Objetivo: o presente estudo foi delineado para verificar se protótipos de mini-implantes (PMI) desenvolvidos para a ancoragem esquelética do aparelho de Herbst são capazes de suportar forças ortopédicas e, também, determinar a variação da força de flexão desses PMI de acordo com o local de inserção (maxila ou mandíbula). **Métodos:** após o cálculo do tamanho da amostra, 13 PMI foram colocados em três cadáveres de Minipigs br1 (seis na maxila e sete na mandíbula). Os corpos de prova foram preparados e submetidos a um teste mecânico. Cálculos da média e o do desvio-padrão foram realizados para cada região. O teste *t* de Student para duas amostras não pareadas foi utilizado para comparar a resistência dos PMI entre as regiões de inserção. O teste *t* de Student para uma amostra foi realizado para o teste de hipótese. Foram consideradas forças ortopédicas aquelas acima de 1,0kgf. **Resultados:** os PMI foram capazes de suportar forças de flexão maiores que 1,0kgf ($13,8 \pm 2,3$ Kg na região posterior da maxila, e $20,5 \pm 5,2$ Kg na região anterior da mandíbula), apresentando significativa menor força de flexão na região anterior da mandíbula ($p < 0,05$). **Conclusão:** os PMI são capazes de suportar forças ortopédicas, sendo mais resistentes quando utilizados na região anterior da mandíbula do que na posterior da maxila, em cadáveres de Minipigs br1.

Palavras-chave: Aparelhos ortopédicos. Implantes dentários. Aparelhos ortodônticos. Procedimentos de ancoragem ortodôntica. Má oclusão de Angle Classe II.

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» The author Klaus Barretto Lopes asserts to be the developer and patent holder of the mini-implant prototypes used in this study.

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INTRODUCTION

Implants and mini-implants have been used as orthodontic anchorages for different purposes in different locations.^{1,2,3} Some researchers have suggested the use of mini-implants as orthopedic anchors in animals^{4,5} and in the treatment of Class III malocclusions with retrusive maxillae in humans.⁶ However, there is little information about the use of mini-implants as orthopedic anchorage in the treatment of Class II malocclusions.

The Herbst appliance has often been used in the treatment of Class II malocclusions, because of its efficiency⁷ and also because of positive effects in orthodontic and orthopedic correction.⁸ Nevertheless, some investigators have stated that the correction of Class II malocclusion is a result of anchorage loss, and could be responsible for negative effects, such as protrusion and gingival recession,⁹ on lower incisors.

Many attempts have been made to reduce the negative effects caused by the Herbst appliance on lower incisors, such as increasing the number of teeth in the mandibular anchorage, using soft-tissue anchorage, splints, and cast splints anchorage.^{10,11} However, these attempts were unsuccessful.

With the intention of solving these problems, a mini-implant prototype was developed for Herbst appliance anchorage, and its flexural resistance was measured in an *in vitro* study.¹² However, a question arose with respect to the resistance strength of this mini-implant prototype when inserted into the bone. The present study was designed to evaluate if the mini-implant prototype developed for Herbst appliance anchorage is capable of withstanding orthopedic forces in Minipigs br1, and to compare the prototype resistance between the sites of insertion.

MATERIAL AND METHODS

Thirteen mini-implants (Neodent, Curitiba, Brazil), 2 mm in diameter and 10 mm in length, with attachment to Herbst appliance telescopic tubes, were inserted in three Minipigs br1^{13,14} (15-month old) after they had been euthanized.

A calculation of the sample size¹⁵ was carried out by means of two pilot studies, one for the posterior region of the maxilla and another for the anterior region of the mandible of one minipig.

Based on the performance of the specimens on the graph, the sample size was calculated with the values ob-

tained with a dislocation of 1.2 mm, because after this point the strength values increased abruptly, indicating the resistance of the metal block.

Afterwards, the following statistical formula was used:

$$n = \frac{(Z \times C_v)^2}{E_r^2}$$

Where n = number of specimens, Z = Standard deviation from normal distribution, C_v = Coefficient of variation, and E_r = Relative error.

After sample size calculation, the number of specimens needed for the final study was 6 in the maxilla and 7 in the mandible.

In order to test sample normality, the Kolmogorov-Smirnov test was carried out. To test the hypothesis, a one-way Student's t test was performed with Minitab 15 (State College, PA, USA). Orthopedic forces above 1 Kgf were considered.^{16,17}

Thereafter, the hypotheses of withstanding orthopedic forces were defined (H₀: μ = 1.0 and H₁: μ > 1.0).

The criterion to reject the null hypothesis was:

$$t_{cal} > t_{\alpha, n-1},$$

where α = 0.05 or calculated:

$$t > t \text{ from } t \text{ table.}$$

To compare the flexural strength between the posterior region of the maxilla and the anterior region of the mandible, a two-way Student's t test was calculated using Minitab 15.

Experiment

The animals were euthanized before the experiment and frozen at a temperature of - 20°C.

Two minipigs received four mini-implants (two in each maxilla and two in each mandible), and one minipig received five mini-implants (two in the maxilla and three in the mandible).

Six specimens with the new mini-implant were obtained from the posterior region of the maxilla, and seven specimens were obtained from the anterior region of the mandible, which were the possible places where the mini-implants could be used to anchor the Herbst appliance.

After that, the straight telescopic tube was placed in the head of the mini-implant with a suitable screw (Fig 1).

To create a guide for the mini-implant, a drill with torque control with a 1.3 mm diameter bur (Neodent, Curitiba, Brazil) was used. The insertion sites were in

the posterior region of the maxilla, between the roots of the upper first molar and the anterior region of the mandible, between the roots of the second premolar and the roots of the third premolar.

To insert the mini-implant, a torque key with a torque meter calibrated at a measure not greater than 30 Kgf.cm was used. When the insertion was completed, new radiographs were taken to check the final positioning of the mini-implants.

To prepare the specimens, the maxilla and the mandible were sectioned into small pieces, using an electric cutting machine. Metal blocks were used to protect the specimens during the experiment. Afterwards, an acrylic resin was used to fix the bone fragments with the mini-implants (Fig 2).

The specimens were placed in an Instron 4400R test machine (Instron, Norwood, MO, USA), with the metal block in the lower part and the telescopic tube in the upper part (Fig 3).

The specimens were submitted to a single cantilever flexure test. Traction was applied at 0.5 mm per minute until 1.5 mm of dislocation was obtained. This value was based on a pilot study carried out by Brettin et al.¹⁸ The values were recorded, and a graph of strength x dislocation was constructed.

RESULTS

Single cantilever flexure tests were successfully carried out on 13 specimens. The graphs illustrate the performance of the specimens during the tests in the maxilla and the mandible (Figs 4 and 5). The mini-implant prototypes showed a flexural strength of 13.86 ± 2.30 Kgf for the posterior region of the maxilla, and 20.5 ± 5.20 Kgf for the anterior region of the mandible.

The normality of the two samples was confirmed with the Kolmogorov-Smirnov test.

For the hypothesis test, the calculated t was 13.71 ($P < 0.001$) for the posterior region of the maxilla. The critical t obtained from the table was 2.015. The criterion used to reject the null hypothesis revealed that:

$$t_{cal} > t_{\alpha, n-1}, \text{ where } \alpha = 0.05 \rightarrow 13.71 > 2.015$$

Therefore, the value calculated for t for the posterior region of the maxilla is outside the region of H_0 acceptance. The null hypothesis was rejected, and the hypothesis that mini-implants cannot withstand orthopedic forces could not be confirmed for the posterior region of the maxilla.

Using the same criterion for the anterior region of the mandible, the calculated t was 9.94 ($P < 0.001$). The critical t obtained from the table was 1.943.

The criterion used to reject the null hypothesis revealed that:

$$t_{cal} > t_{\alpha, n-1}, \text{ where } \alpha = 0.05 \rightarrow 9.94 > 1.943$$

Similarly to the maxilla, the value calculated for t is outside the region of H_0 acceptance. The null hypothesis was rejected, and the hypothesis that mini-implants cannot withstand orthopedic forces could not be confirmed for the anterior region of the mandible.

A statistically significant difference was found in the flexural strength between the posterior region of the maxilla and the anterior region of the mandible ($P = 0.015$): the anterior region of the mandible was significantly more resistant.

DISCUSSION

One of the purposes of this study was to quantify the flexural resistance of the mini-implant prototypes when inserted in the posterior region of the



Figure 1 - Straight telescopic tube placed in the head of the mini-implant with a suitable screw.

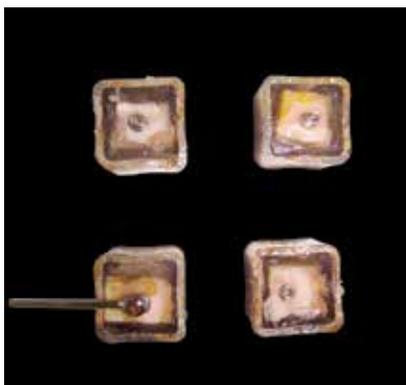


Figure 2 - Bone fragments with mini-implants inserted which were included in the metal block.

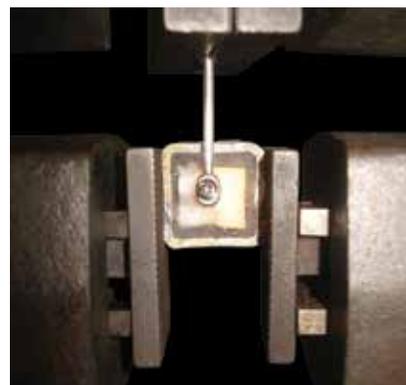


Figure 3 - Specimen ready for flexure test.

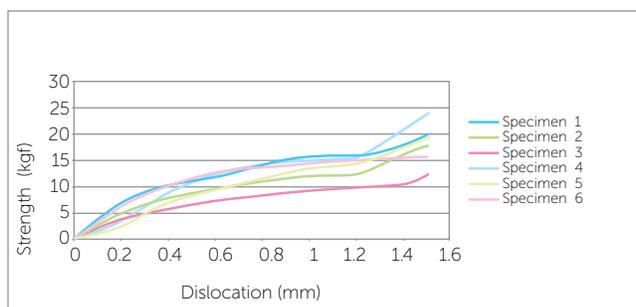


Figure 4 - Graph of strength x dislocation of six specimens tested in the posterior region of the maxilla.

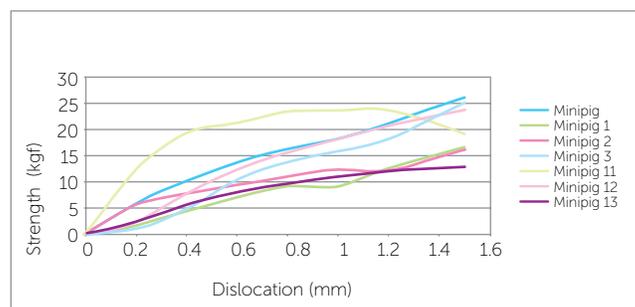


Figure 5 - Graph of strength x dislocation of the seven specimens tested in the anterior region of the mandible.

maxilla and the anterior region of the mandible. Although studies involving extrapolations from animals to humans should be viewed with caution, the authors' intention was to assess the strength that the mini-implant prototypes could withstand in specific regions in a Minipig br1. The mini-implant prototype showed a flexural strength of 13.86 ± 23.0 Kgf for the posterior region of the maxilla, and 20.55 ± 5.20 Kgf for the anterior region of the mandible. Another purpose of this study was to test the hypothesis that mini-implant prototypes are not capable of withstanding orthopedic forces. The hypothesis was rejected.

Miyawaki et al³ reported that the success rate of mini-implants with a 1.0-mm diameter was significantly lower than that of other mini-implants with 1.5-mm or 2.3-mm diameters. This author suggested the use of mini-implants with a 1.5-mm diameter for patients with average-to-low mandibular plane angle, and a 2.3-mm diameter for patients with a high mandibular plane angle (i.e., with a thin cortical bone). Miyawaki et al³ did not find a significant association between the success rate and the mini-implant length. However, Brettin et al¹⁸ concluded that bicortical mini-implants provide superior anchorage resistance, reduced cortical bone stress, and superior stability when compared with monocortical mini-implants. Also, according to Barros et al¹⁹ increases in mini-implant diameters significantly influenced the increases in placement torque and fracture torque on quantities that progressively reduced the fracture risk. Therefore, in order to increase the resistance of the mini-implant prototype, the diameter was increased to 2.0 mm and the length to 10 mm, so as to achieve bicortical anchorage.

It could be argued that the 2.0-mm diameter mini-implant is too large for the inter-radicular space. However, Poggio et al.²⁰ showed that, in humans, this size is compatible with the inter-radicular distance between the upper first molars and second premolars and between the lower cuspids and the first premolars (3 mm when the distance from the alveolar crest is 8 mm). This is the location suggested for the insertion of the mini-implant prototypes for the Herbst appliance anchorage.

Two pilot studies were performed before the experiment to test the researchers' abilities and to calculate the sample size.¹⁵ The number of specimens was calculated in order to obtain scientific validation with the fewest possible specimens and animals.

In the present study, the mini-implant prototypes were inserted after the animals had been euthanized. Huja et al²¹ explained that no healing period or adaptive response could occur, and resistance strength was an indication of primary stability. Therefore, the values found in the present study are similar to the experiments of immediate loading after placement. Some studies have evaluated bone contact according to the healing period, recommending immediate loading after placement.^{1,22,23}

Storage conditions of the specimens used in the study have been associated with differences in pull-out strengths during mechanical traction testing. Simonian et al²⁴ found lower pull-out strengths in frozen specimens. However, Roe et al²⁵ reported no significant difference when the test was performed no later than one week after storage at -20° C. Therefore, to avoid any influence of storage conditions on the results, the specimens used in this study were prepared

on the same day or on the day after insertion of mini-implants, frozen at -20°C and tested between one and seven days after mini-implant insertion.

A single cantilever flexure test was performed instead of a shear force test and traction test, because there was a distance of 4 mm between the block base and the point of force application. In a shear force test, the point of force application should be parallel to the block base. A traction test would not reproduce the perpendicular strength received by the mini-implant when used as an anchorage for the Herbst appliance.

Brettin et al¹⁸ also performed a cantilever flexure tests, but in human cadavers. They found lower values than those of the present study (20.55 kgf for bicortical mandibular mini-implants and 13.86 kgf for monocortical maxillary mini-implants in Minipigs, against 11.0 kgf for bicortical mandibular mini-implants and 9.0 kgf for bicortical maxillary mini-implants, 5.0 kgf for monocortical maxillary mini-implants and 7.0 kgf for monocortical mandibular mini-implants in human cadavers). This difference may be related to the larger diameter of the mini-implants used in the Minipigs (2.0 mm in the Minipigs against 1.6 mm in the human cadavers) or increased cortical bone thickness in the Minipigs.

The cantilever test is a static test. Probably, if dynamic forces had been applied to the mini-implants, different results would have been obtained. Future *in vivo* studies using the Herbst appliance anchored in bone will answer this question.

According to Huja et al²¹ some possible problems were related to cantilever tests, namely: standardization, reproducibility, bone flexure and collision of the mini-implants with adjacent root of the tooth. These problems could negatively influence accuracy of results regarding the flexural strength of the mini-implants. In this research, these possible problems were avoided, because the flexural strength of the mini-implants was obtained following the experiment performed by Brettin et al¹⁸ in which the mini-implants were tested with dislocation of 1.5 mm. Those authors performed a pilot study in cadavers in which they concluded that the mini-implants could present mobility with dislocation above 1.5 mm.

To test the hypothesis, orthopedic forces above 1.0 kgf were considered. This value is very common in Orthodontics, in clinical work and research,

because of the headgear which functions with 500 g per side.^{16,17} Other studies showed the use of orthopedic forces (from 500 to 800 g) on implants^{4,5,6} and a few focused on the use of orthopedic forces in mini-implants, such as the study by Büchter et al²⁶ who found that mini-implants resisted up to 900 g without mobility.

The flexural strength differed significantly between the posterior region of the maxilla and the anterior region of the mandible. However, this result may have been influenced by the fact that the mini-implants fixed in the mandible had bicortical anchorage while those fixed in the maxilla apparently did not have this anchorage. Probably, the distance between the cortical bones in the maxilla in the posterior region was greater than 6 mm or the cortical-lingual was not accessible due to the palate being too low. Because the mini-implant was 10 mm long, the active part was only 6 mm. This suggests that in future studies the mini-implants should be constructed according to the distance between the cortical bone, as Brettin et al¹⁸ did with human cadavers.

Other forces that could affect the resistance of the mini-implant prototypes could be those originating from muscles and soft tissues of the face, due to mandibular advancement caused by functional appliances. However, earlier studies^{27,28} showed that the forces delivered to the teeth by functional appliances were of low intensity (80 and 160 gf).

The question that arises is whether to use mastication forces as a parameter rather than the orthopedic forces. According to Pancherz and Anehus-Pancherz²⁹, there is no contact in the posterior teeth after the installation of the Herbst appliance. The contact occurs only in the anterior teeth. This is responsible for the decrease in mastication efficiency as well as in temporal and masseter muscle activity in the first three months of treatment. After that, the authors observed an increase in the mastication forces during six months of treatment.

The forces transmitted to the teeth when using the Herbst appliance are probably the best parameter. However, it was not possible to locate studies with the data that is necessary to carry out a statistical analysis, and the force transmitted to the mini-implants in the skeletal anchorage of the Herbst appliance remains unknown.

Thus, *in vivo* and clinical studies are necessary to assess the possibility of using the Herbst appliance skeletal anchorage in humans.

CONCLUSION

The mini-implant prototypes developed for Herbst appliance skeletal anchorage are capable of withstanding orthopedic forces in Minipigs br1 cadavers. The prototypes are more resistant in the anterior region of the mandible than in the posterior region of the maxilla.

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