

# Morse taper internal connection implants: would abutment reseating influence retention?

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## Abstract

**Aim:** To investigate whether the removal and reseating of the abutment influences the retention of Morse taper implant system. **Methods:** Two Morse taper implant systems were selected: Bicon Dental Implant™ system (Bicon, LLC, Boston, MA, USA) and Kopp Implant® system (Kopp®, Curitiba, PR, Brazil). In both systems, the abutment is connected to the implant with a locking taper. To seat the abutment, the same vertical force was applied in both systems. It was measured the compressive force necessary to engage effectively the locking taper connection and the tensile force to displace it. **Results:** The compressive force was determined by four activations in each abutment-implant set, and the sum of these forces was 21 N and 17 N in the Bicon™ and Kopp® systems, respectively. Next, a tensile test was performed, revealing that the Bicon™ system presented a 208 N resistance, whereas it was 194 N in the Kopp® system. Other three rounds of compression and tensile loads were applied, removing and reseating the abutment. We obtained the following tensile values: 367 N, 500 N and 756 N in the Bicon™ System and 336 N, 360 N and 420 N in the Kopp® system. **Conclusions:** When the sets were subjected to repeated rounds of compressive and tensile forces, displacing and reseating the abutment, the tensile value increased.

**Keywords:** dental implant-abutment design; dental implants; compressive strength; tensile strength.

## Introduction

The search for an optimal implant configuration has led to various internal geometries with different prosthetic platforms<sup>1</sup>. The goal is to reduce the fatigue on the setting screws and obtain a better biological seal, mechanical strength and aesthetics<sup>2-5</sup>.

One of the main challenges confronting implant dentistry today is to achieve an implant-abutment connection that satisfactorily meets the principles of biomechanics. This difficulty is often due to loosening or even breaking of the fixation screw<sup>6-8</sup>. Screwless implant systems have several advantages, such as the possibility of appropriate cervical region aesthetics and fewer prosthetic components. It leads to cost reduction, simpler clinical procedures, and greater fracture resistance of the prosthetic component<sup>3-4,9-10</sup>. Moreover, the Morse cone design platform offers a greater and stronger implant-abutment interface, yielding a better fit between these elements. It allows gap reduction and increased micro-movement resistance. It also provides joint stability<sup>5</sup> due to the high frictional force created by the locking taper, which is produced by the pressure of two

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sliding surfaces. As a result, surface oxide layers break down, a phenomenon known as cold welding<sup>11</sup>. Therefore, there is less risk of gaps between the implant and abutment, producing a seal proven to delay or even prevent bacterial invasion<sup>12-14</sup> and to be clinically reliable<sup>15</sup>. A recent study on implant components stability demonstrated that loosening in sets with internal tapered connections is less frequent than in sets with other connection types<sup>16</sup>.

Although high success rate in Morse taper implant systems has been reported<sup>3-4,9,15,17-18</sup>, implant treatment is often hindered by complications<sup>1,19</sup>. Abutment displacement may occur<sup>9,17</sup>. Furthermore, removing the abutment-crown complex may be required. This raises the question of whether the reinsertion procedure will jeopardize the system retention. Additionally, doubts were raised regarding quantification of forces applied during installation of frictional systems<sup>19</sup>. It should be noted that in Morse taper systems, the manufacturer's recommendation is to insert, align and then put the abutment in place by seating taps intermittently applied using a weight load. This load is applied to achieve the required mechanical friction between the external wall of the abutment and the internal wall of the implant, which results in high contact pressure<sup>1</sup>. In screw-retained systems, the screw is tightened with a torque established by the manufacturer<sup>19</sup>. To the best of our knowledge, in the Morse taper system there is no recommendation as to the compression required to obtain optimal imbrication. Given the aforementioned considerations, the following questions arise: a) Will the intensity of the applied vertical force modify the tensile strength? b) Will the removal and repositioning of the abutments influence their retention? The present study addresses the latter.

## Material and methods

In this study, two commercial brands of Morse taper implant systems, Kopp® (Kopp®, Curitiba, PR, Brazil) (Figure 1) and Bicon™ (Bicon Inc., Boston, MA, USA), were analyzed, both with implant-abutment sets accomplished by mechanical and frictional imbrication. They have a 1.23-1.5 degree tapered post that fits into a mirror-image shaft, with no screw.

An implant and an F II straight abutment model were selected from the Kopp® company measuring 4.3 x 13 mm and 4.5 x 13 mm, respectively. Similarly, an implant and abutment set was selected from the Bicon™ company measuring 4.0 x 11 mm and 4.0 x 6.5 mm, respectively (Figure 1). For each set, a rigid base was fabricated consisting of a bolt (M 12 x 1.25 mm) with a hole at the top drilled with a mechanical lathe with the appropriate diameter to fit each implant tightly. To obtain the greatest possible resistance, a metal anaerobic adhesive (Three Bond 1375 - ThreeBond of Brazil Ltda) was used.

The sets were fixed individually in a Universal Testing Machine (DL 500 Emic - Equipment and Testing Systems Ltd., PR, Brazil). Then, they were connected to a load cell (Model CCE5KN - Emic) with a capacity of 5000 N at the machine's lower section. A weight (BCP, Kopp®, Curitiba,

PR, Brazil) was used to generate compressive load. This device consisted of a constant-length (0.065 m) metal rod weighing 0.188 kg. It was set in the upper part of the machine in a position concentric to the impact body trajectory to promote the attachment of the abutment to the implant (Figure 2).

In the first phase of the experiment, the abutment-implant sets were activated four times each by means of compressive force. Briefly, a single operator positioned the abutment over the implant without any pressure. Next, engagement between the complementary tapered surfaces of implant and abutment was accomplished by the impact of the body weight, dropped from a known height ( $h=0.065$  m), limited by the instrument. The force was applied along the long axis of the implant.

In the second phase of the experiment, the abutment of each set was fixed to a forceps shaped device connected to the same load cell, but now in a reversed position, on the upper part of the Universal Testing Machine (Figure 3). Next, the sets were subjected to a tensile force in the same machine until the abutment-implant assembly was displaced. The tensile force values required for the removal of abutment were recorded.

In the last phase of the experiment, the abutments were removed and resealed again three times, summing four reseating/removal cycles. At each displacement of the abutment, the tensile force necessary to remove was recorded, enabling an evaluation of whether the applied tensile force was greater than, equal to, or less than that obtained the first time. Thus, it was checked whether the removal and repositioning of the abutments increased the friction and therefore the retention values, or if there was a friction loss and consequent loss of retention.

## Results

In the first phase of the experiment, the compressive load applied by the weight (0.188 kg) on the Kopp® and Bicon™ abutment-implant sets was measured by the load cells. The experiment was performed under identical conditions for both brands. There were four activations in



Fig. 1: Kopp® and Bicon™ implant-abutment sets fixed to a rigid base.

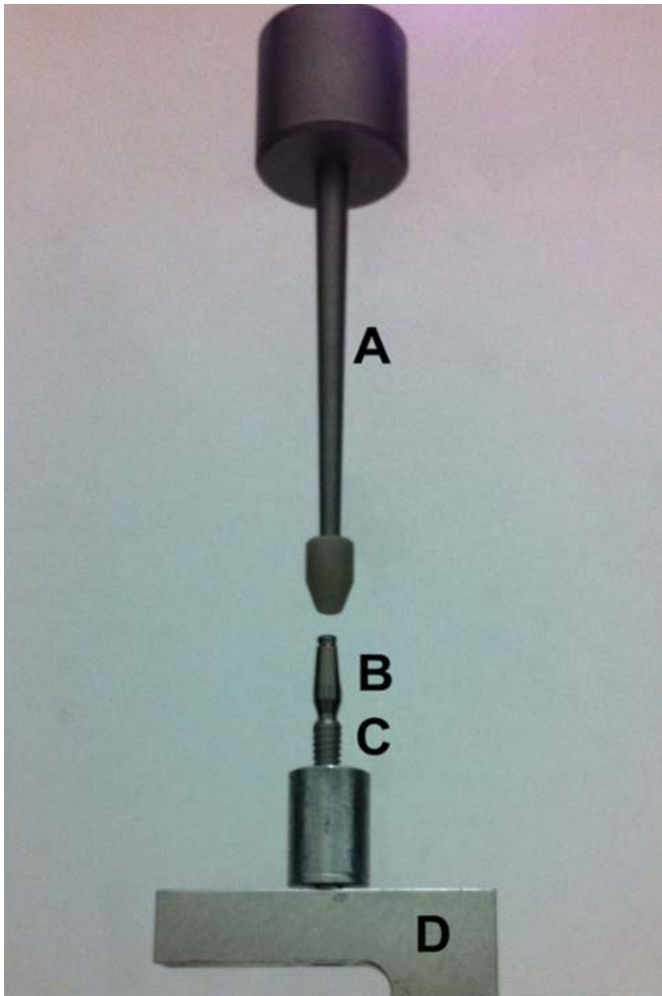


Fig. 2: Implant-abutment set positioned in the Universal Testing Machine for compressive test. A: weight in position; B: abutment; C: Implant; D: Load cell.

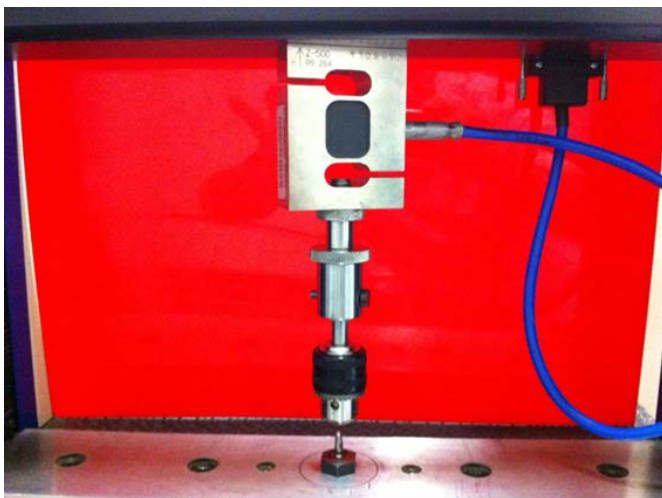


Fig. 3: Implant-abutment set positioned in the Universal Testing Machine for tensile test.

each set and the total results were 17 N for the Kopp® set and 21 N for the Bicon™ set.

In the second phase of the experiment, tensile loads were applied under identical conditions on the two implant-

abutment sets. It was observed that the Bicon™ system presented a 208 N resistance and the Kopp® system, 194 N.

In the last experiment, when new compressive and tensile loads were re-applied three times, it was possible to measure the following tensile values: 367 N, 500 N and 756 N for the Bicon™ system and 336 N, 360 N and 420 N for the Kopp® system.

## Discussion

This study aimed to investigate whether the removal and reseating of the abutment influences the Morse taper implant system retention. High success rate in Morse taper implant systems has been reported<sup>9,15,17-18</sup>. However, an implant treatment may be hampered by complications, abutment displacement may occur<sup>9</sup> and removal of the abutment may be necessary. This raises the question of whether the reinsertion procedure will risk the system retention<sup>17</sup>.

The present study showed that the tensile strength was proportional to the seating force applied to the abutment. In the Kopp® system implant-abutment set, in which the impact force was lower (17 N), the force to remove the abutment was 194 N. In the Bicon™ set, with a greater seating force (21 N), the force required to pull out the abutment was 208 N, in the same conditions. These findings are in agreement with the statement that locking taper systems, when inserted into the implant, require a removal force greater than the insertion force<sup>9</sup>.

In the final test, when the abutments were removed and then inserted three more times, tensile strength values of 367 N, 500 N and 756 N were obtained with the Bicon™ system and 336 N, 360 N and 420 N with the Kopp® system. According to these results, Bicon™ and Kopp® implant systems, when subjected to the same compressive loading conditions, had similar behavior. It means that the greater the applied compression, the greater is the tensile strength, although the obtained values were different.

Previous research has shown a positive correlation between the number of tappings and the removal forces for a locking taper implant system<sup>9</sup>, which is consistent with our findings. In fact, these results may be explained by the phenomenon of superficial hardening, which is the increased induration of the outer and inner surfaces of the implant and abutment, varying according to the elasticity of each metal alloy. With greater impact force there is a tendency of the abutment to undergo superficial plastic deformation on its platform (Figure 4a).

Additionally, our results can be explained by the phenomenon of the buckling of the abutment. When compressing a rod with a large diameter, a slight inclination along the axis is produced (Figure 4b). Finally, these results can also be explained by the fact that the tension between the surfaces tends to be asymptotic, due to their angulation, which creates a non-uniform distribution of the compressive stresses and consequently changes the cross-sectional rigidity (Figure 4c). Therefore, the tensile strength should increase as the compressive force increases. When these phenomena

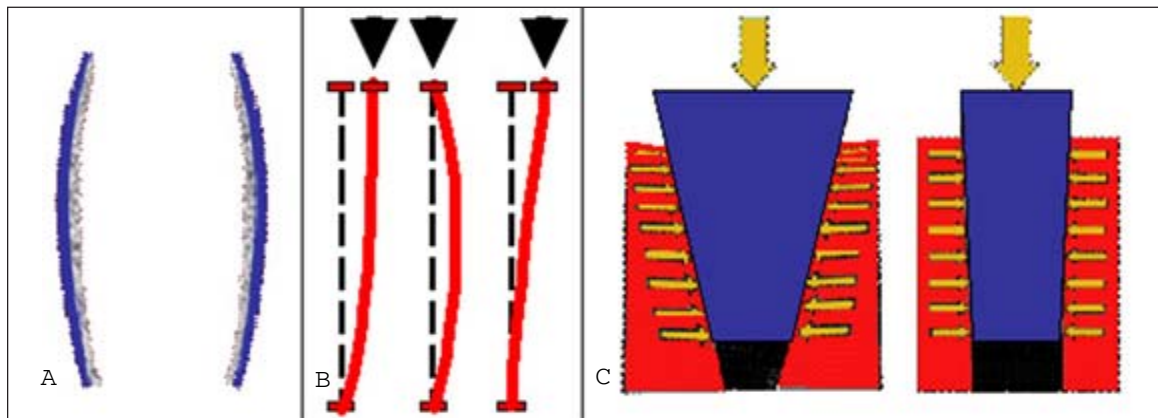


Fig. 4: Schematic drawings: a) superficial hardening; b) buckling; c) asymptotic tension superficial induration.

occur, the individual compressive force may not be concentric, generating a variable compressive load. Although this increase in surface rigidity could be interesting, cracks may occur, which is an undesirable effect<sup>20</sup>.

It should be mentioned that *in vitro* study models might not predict the clinical reality. There are many factors influencing *in vivo* implant components placement and behavior. As a matter of fact, this *in vitro* study did not consider bending forces, usual in a clinical situation, which may alter the abutment resistance to displacement. Furthermore, to the extent of our knowledge only the two systems analyzed in this work use a tapered interference fit solely. Thus, it is difficult to compare our data with other researches. In a recent systematic review about conical abutment connection implants, the majority of the Morse taper implant system designs included a screw<sup>4</sup>. Nonetheless, this experimental work sheds light on the influence of abutment reseating on the retention of Morse taper implant systems. Although only one sample of each implant system was used, the mechanical tests revealed a clear tendency with important practical application.

In the present study, we observed that higher compressive forces generated higher tensile strength. The results obtained in this study reveal that the removal of the abutment and subsequent repositioning for a total of four repetitions increased the tensile force required to remove the abutment, thereby increasing its frictional retention. Within the limits of this *in vitro* study, the reseating of abutments in locking-taper implants seems to be a successful procedure. Further research is necessary to clarify the clinical relevance of our findings, and to establish the optimum compressive force for each commercial brand that uses exclusively the Morse taper fixation system to connect implant and abutment.

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