

Stability Evaluation of Micro-Screw Implant in Cases Given a Bone Inducing Substances "An Experimental in Vivo Study"

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الخلاصة

أهداف: لتحديد الوقت الأمثل لتحليل الغرسات التقويمية وذلك بواسطة تقييم الثباتية وكذلك كشف إمكانية استخدام الوندرونات الصوديوم كعقار طبي لمنع ذوبان العظم وبالتالي إمكانية زيادة ثباتية الغرسة التقويمية. **المواد وطرائق العمل:** ثمانية وأربعون غرسة تقويمية، اثنا عشر أرنب حقلية وأربعة عشر أمبولة من الوندرونات الصوديوم. قسمت الأرناب إلى مجموعتين رئيسيتين، مجموعة خضعت لعلاج وأخرى بدون علاج للمقارنة، قسمت كل مجموعة إلى ثلاث مجاميع فرعية: التحميل الآني للقوة بعد الغرس، تحميل القوة بعد أسبوعين من الغرس والأخيرة هي تحميل القوة بعد أربع أسابيع من الغرس ومن ثم تم قياس الثباتية باستخدام جهاز الثباتية البريوتيسست لجميع المجاميع الفرعية بعد الغرس وقبل التحميل وبعد التحميل وبعد أسبوعين من التحميل. **النتائج:** أظهرت نتائج الدراسة عدم وجود اختلاف معنوي في المجاميع الفرعية المعرضة للعقار قبل تحميل القوة وبعد أسبوعين من تحميل القوة لكن الاختلاف كان معنوي بعد تحميل القوة مباشرة، كذلك الحال بالنسبة للمجموعة الفرعية المحملة أنيا لا يوجد اختلاف حتى بعد أسبوعين من التحميل. أما فيما يخص المجموعة الفرعية المحملة بعد أسبوعين وأربعة أسابيع فقد أظهرت نتائج اختلاف معنوي في جميع قراءات مع ملاحظة إن الثباتية الآنية هي الأفضل تقريبا. **الاستنتاجات:** من نتائج البحث يمكن القول ان الغرسات التقويمية حتى الأصغر حجما يمكن استخدامها كمثبت جيد مع جهاز التقويم مع إمكانية تحميلها مباشرة بعد الغرس، كذلك وقد أظهرت النتائج إن استخدام عقار الوندرونات صوديوم لايدر فائدة على ثباتية الغرسة التقويمية على غرار المعايير المستخدمة في هذه الدراسة.

ABSTRACT

Aims: The present study aimed to investigate the desirable loading time of micro-screw implant by stability evaluation and detecting the possibility of using alendronate sodium to increase the stability. **Materials and methods:** Forty eight micro-screw implant, twelve adult rabbits and fourteen alendronate sodium ampoules were used in this study, the rabbits were divided into two main groups, treated and control group, which further subdivided into three subgroups. Eight micro-screw implant instilled in tibiae of each subgroup just six of them used for test the remaining cancelled, the stability test down using the periotest. These subgroups were nominated according to the loading times which are immediate loading, loading after two weeks and loading after four weeks with stability measured after instillation, before and after loading and two weeks after loading. **Results:** No significant differences between subgroups before loading and after two weeks of loading but significant after loading. For immediately loaded treated subgroup no significant differences between immediate loading and two weeks after loading. For two and four weeks treated subgroup a significant difference in stability between immediate instillation and after loading. **Conclusions:** Micro-screw implant even smaller diameter could be used as a fixed anchorage in orthodontics and possibly could be loaded safely from time of immediate instillation, further the use of alendronate sodium add no benefit to increase stability according to the criteria used in this study.

Key words: Stability, Micro-screw implant, Bone inducing substances.

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INTRODUCTION

Good anchorage control is one of the prerequisites of successful orthodontic therapy. Recently, micro-screw implant have been proven to be useful in establishing absolute anchorage without the use of extra-oral appliances, add to their flexibil-

ity in choosing implant locations, lower medical costs, simpler implant surgery, and lower degrees of discomfort after implantation compared with traditional dental implants.^(1,2) However, these mini-screws loosen easily and their failure rate may be as high as 25%.^(3, 4)

The stability is of two components: *Primary stability* which is established from the mechanical lock between the micro-screw implant surface and bone.⁽⁵⁾ It is depends on the thickness and integrity of the cortical bone. The Micro-screw implant design, and loading protocol.^(5,6) *Secondary stability* is achieved through continuous bone remodeling around the Micro-screw implant, leading to osseointegration (increasing bone density).^(7,8) Which is a critical determinative factor of the performance of endosseous implant.^(9,10) Healing time, has a considerable impact on the osseointegration.⁽¹¹⁻¹⁴⁾

Some studies showed increased bone marrow density (BMD) of 4% to 8% with the use of alendronate sodium, Bone-resorption inhibitor. Osteoclast inhibition is the primary reason for this phenomenon.⁽¹⁵⁾ There are no studies comparing the effect of systemic administration of alendronate on dental implant osseointegration. In this study, a rabbit tibia model was used to examine the effects of alendronate as osteoclast inhibitors at different loading and healing times on mechanical stability. The present study aimed to investigate the desirable loading time of micro-screws by stability evaluation.

MATERIALS AND METHODS

Fourty eight commercially available self-drilling titanium micro-screw implants (Abso-anchor, Dentos Inc, Daegu, Korea), twelve adult female rabbits and fourteen alendronate sodium ampules (Diamond Pharma-Damascus-Syria under license of ABC Farmaceutici-Torino-Italy) for intra muscular injection were used in this study. Each micro-screw implant was selected from the same series, with the same length, diameter and similar shape. measuring 1.3 mm in diameter and 5 mm in

length, the rabbits were 12 months old (each weighting 2 kg) and the drug Pharmacologic class; Bisphosphonate, Therapeutic class: Bone-resorption inhibitor, its Action is impeding bone resorption by inhibiting osteoclastic activity, absorbing calcium phosphate crystal in bone, and directly blocking dissolution of hydroxyl apatite crystal of bone.

All micro-screws and rabbits were divided into two groups; the first untreated group (control group) is the plane group (p) which further subdivided into three subgroups, which are, the immediate loading plane subgroup (0WP), 2-week healing plane subgroup (2WP) and 4-week healing plane subgroup (4WP) while the second group is the treated group (T) which also subdivided into three subgroups, the immediate load treated subgroup(0WT), 2-week healing treated subgroup (2WT) and 4 week treated subgroup (4WT). The drugs given to each rabbit intra muscularly (0.3 ml) for fourteen days before Micro-screw implant instillation (which is just one of the multiple courses recommended, to human, by the manufacturer). There were 8 micro-screws in each subgroup six of them tested and two cancelled due to failure like fracture of screw during driving and some shows bone crack at penetration of screw. There were two rabbit in each of the six subgroups. These six subgroups may encounter the three pathological periods of bone healing after the instillation of the micro-screw implant, that is, the traumatic period, granulation period, and callus period, respectively.⁽¹⁶⁾ The micro-screw implant were instilled in the tibiae of each animal, all the microscrew implant were then tested for their stability using the periotest machine (Medizintechnik Gulden e.k. Eschenweg 3, 64397 Modautal, Germany) (Figure 1)

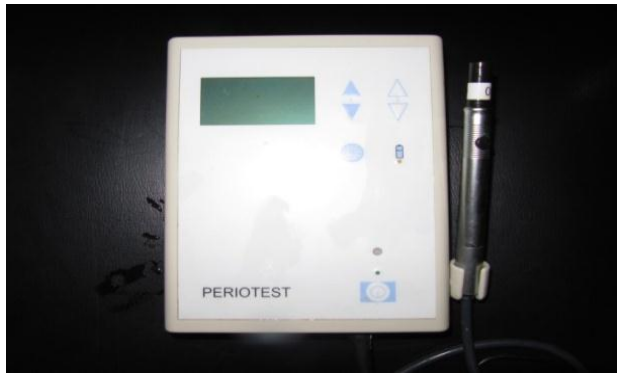


Figure (1): Periotest machine

Which is specified for testing the stability of any structure inserted in the bone and also for natural tooth structure. The periotest scale extend from -8 to +50, the lower the value, the greater is the stability with the periotest value range from -8 to 0 this mean a good osseointegration and implant can be loaded, +1 to +9 values mean clinical examination is required and loading is not yet possible, more than +9 mean that the implant must not be loaded.

Surgical Procedures

All surgeries were performed under sterile conditions in an animal operation room. The animals were anesthetized intramuscularly with a combination of ketamine (44 mg/kg of body weight) and xylazine (7 mg/kg of body weight). The local nerve supplies of the internal surface of the tibia were further blocked with 0.5 ml of 2% Lidocaine (Figure 2).



Figure (2): the local nerve further anaesthetized with 0.5ml 2% lidocaine

The tibiae body was exposed by incisions through the skin, fascia, and periosteum (Figure 3)



Figure (3): incisions through the skin, fascia and periosteum

The cortical bone of the preparation sites was penetrated using a 0.6mm-diameter guide drill under profuse irriga-

tion (Figure 4). After pilot drilling, the Micro-screw implants were placed using a manual driver (Figure 5).



Figure (4): guide drill under perfused irrigation



Figure (5): manual drilling of Micro-screw

All Micro-screw implants were allowed to penetrate the first cortical layer and going through the woven bone only (not penetrating the opposing cortical plate). The loading involved nickel-titanium closed-coil springs (Denturum)

were applied to the coronal portion of the Micro-screw implants with 100g of force using tension gauge (Anthogyr company, France) (Figure 6 a,b). the mucoperiosteum and muscle were sutured in separate layers using absorbable sutures (Figure 7).



Figure (6a): application of coil spring



Figure (6b): application of coil spring

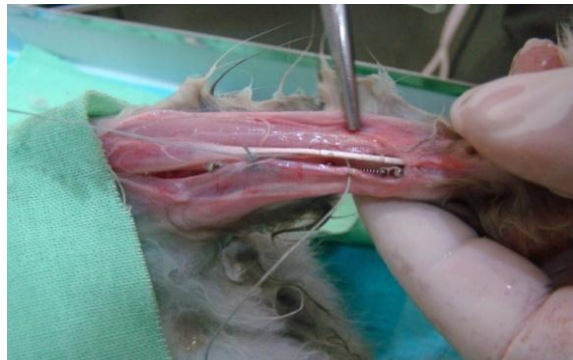


Figure (7): suturing the tissue layers with absorbable suture

The stability testing involve the following: In the 0WP subgroup the implant were tested for their stability immediately after instillation (figure 8) then after loading (figure 9) the test repeated after 2 week healing period. For the 2WP subgroup; the stability tested immediately

after micro-screw implant instillation then the specimen left 2 week for healing then the surgical site opened again. The test repeated again before loading then after loading each rabbit then lifted for healing period of two week which then scarified.



Figure (8): stability test before loading



Figure (9): stability test after loading

The stability again tested. For the 4WP subgroup; the stability tested after instillation of the micro-screw implant then lifted for healing period of 4 week after that the surgical site opened again and the stability tested before loading and after loading then each rabbit lifted for healing period of 2 weeks, which then scarified and the sta-

bility tested again.

RESULTS

The statistical analysis showed differences, but not significant between subgroups 0WT, 2WT and 4WT before loading Table (1), Figure (10).

Table (1): ANOVA statistical test for treated subgroups before loading

	Sum square	Degree of freedom	Main square	F value	P value
Between group	0.048	2	0.024	3.116	0.074
Within group	0.115	15	0.008		
Total	0.163	17			

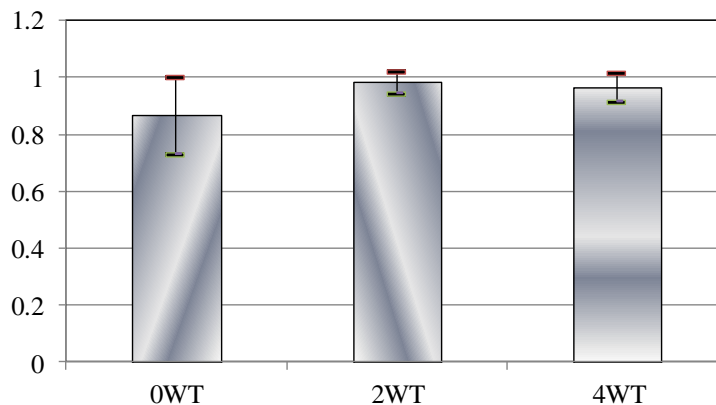


Figure (10): Histogram of Duncan statistical test between treated subgroups before loading.

After loading no significant differences between 2WT and 4WT, but significant for 0WT Table (2), Figure (11) while

on comparing the results between groups after a healing periods of 2 week.

Table (2): ANOVA statistical test for treated subgroups after loading

	Sum square	Degree of freedom	Main square	F value	P value
Between group	0.063	2	0.032	11.400	0.001
Within group	0.042	15	0.003		
Total	0.105	17			

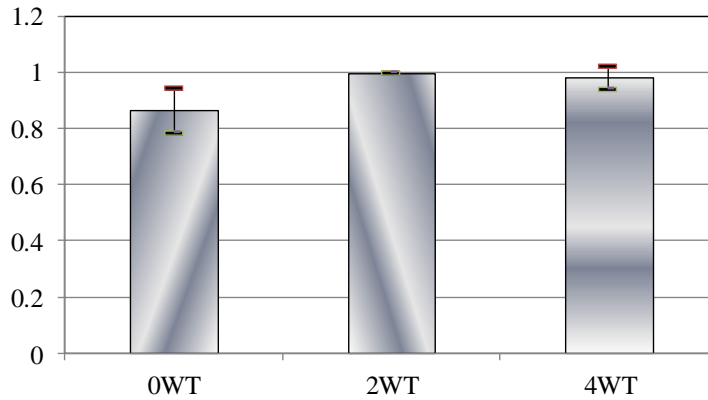


Figure (11): Histogram of Duncan statistical test between treated subgroups after loading

The result showed no significant differences between them Table (3), Figure (12).

Table (3): ANOVA statistical test for treated subgroups after healing period followed loading

	Sum square	Degree of freedom	Main square	F value	P value
Between group	0.043	2	0.022	3.197	0.070
Within group	0.102	15	0.007		
Total	0.142	17			

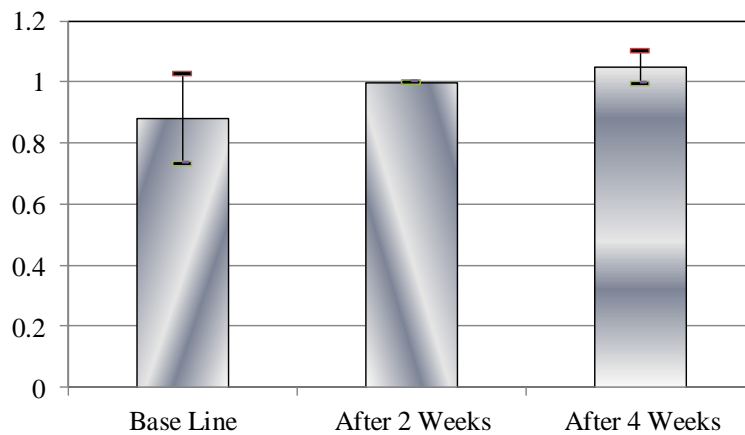


Figure (12): Duncan statistical test for two weeks treated subgroups.

On comparing the results of stability for 0WT subgroup before (BL) and 2 weeks healing after loading (TWAL) also after

immediate loading (IL) and 2 weeks healing after loading no significant differences observed Table (4,5).

Table (4): Statistical T test for stability of unloaded micro-screw implant and two weeks after loading

	Mean	Standard deviation	T test	Degree of freedom	P value
BL	0.8667	0.13663	0.894	10	0.392
TWAL	0.9333	0.12111			

Table (5): Statistical T test for stability of loaded micro-screw implant and two weeks after loading

	Mean	Standard deviation	T test	Degree of freedom	P value
IL	0.8667	0.08165	1.118	10	0.290
TWAL	0.9333	0.12111			

For 2WT subgroup a significant differences had been shown in stability among immediate instillation, loading after two weeks and 2 weeks after loading Table (6) Figure (13), for 4WT subgroup

also significant differences among immediate instillation, loading after 4 weeks and 2 weeks after loading Table (7) Figure (14).

Table (6): ANOVA statistical test of stability differences among immediate instillation, loading after 2 weeks and 2 weeks after loading

	Sum square	Degree of freedom	Main square	F value	P value
Between group	0.088	2	0.044	5.338	0.018
Within group	0.123	15	0.008		
Total	0.211	17			

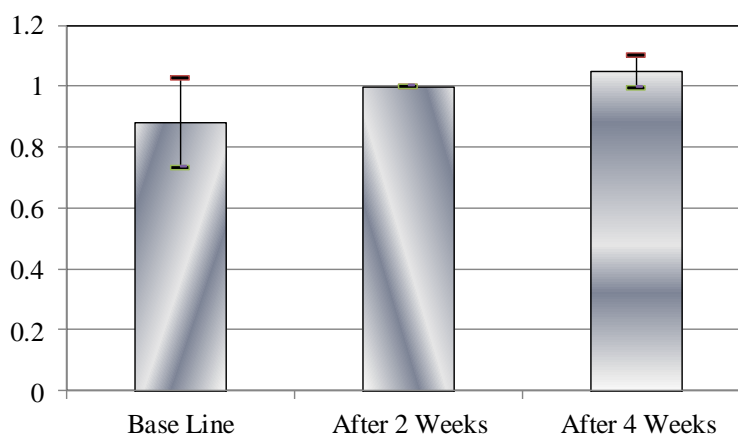


Figure (13): Duncan statistical test for two weeks treated subgroups.

Table (7): ANOVA statistical test of stability differences among immediate instillation, loading after 4 weeks and 2 weeks after loading

	Sum square	Degree of freedom	Main square	F value	P value
Between group	0.063	2	0.032	6.196	0.011
Within group	0.077	15	0.005		
Total	0.140	17			

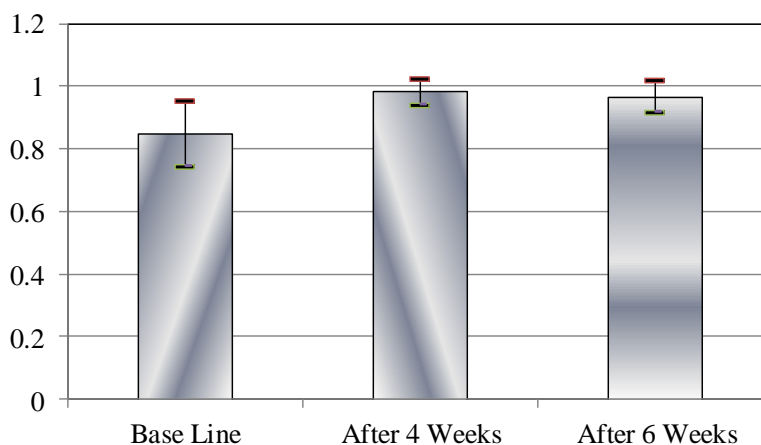


Figure (14): Duncan statistical test for four weeks treated subgroups.

For the plane subgroups, a significant difference of 0WP subgroup from both 2WP and 4WP subgroups which showed no significant differences. This is before and

after loading (Table 8, 9) but after a healing period of two weeks no significant differences observed (Table 10).

Table (8): Duncan statistical test for the control sample of the three subgroups after loading

	Mean	Standard deviation	Duncan
0WP	0.7167	0.16021	A
2WP	1.0	0.0	B
4WP	0.9667	0.05164	B

Table (9): Duncan statistical test for the control sample of the three subgroups before loading

	Mean	Standard deviation	Duncan
0WP	0.60	0.12649	A
2WP	1.0	0.0	B
4WP	0.90	0.06325	B

Table (10): Duncan statistical test for the control sample of the three groups after after loading

	Mean	Standard deviation	Duncan
0WP	0.9167	0.04082	A
2WP	0.9833	0.09832	A
4WP	0.8833	0.07528	A

On comparing the results of stability for 0WP subgroup before loading and after 2 weeks of loading also after immediate

loading and after 2 weeks a significant differences observed Table (11,12)

Table (11): Statistical T test of immediately instilled screw and two weeks after loading

	Mean	Standard deviation	T test	Degree of freedom	P value
BL	0.6	0.12649	5.836	10	0.000*
IL	0.9167	0.04082			

*significant difference

Table (12): Statistical test of loaded screw and two weeks after loading

	Mean	Standard deviation	T test	Degree of freedom	P value
BL	0.7167	0.16021	2.963	10	0.014*
TWAL	0.9167	0.04082			

*significant difference

For 2WP subgroup a significant difference in stability of immediately instilled micro-screw implant from stability of

loading after 2 weeks and those lifted for 2 weeks after loading (Table 13).

Table (13): Duncan statistical test for the two weeks control subgroup

	Mean	Standard deviation	Duncan
0WP	0.8500	0.05477	A
2WP	1.0	0.0	B
4WP	0.9833	0.09832	B

for the 4WP subgroups a nearly significant differences in stability reading in the three times intervals, since time of immediate

instillation, 4 weeks then loading and 2 weeks after loading (Table 14).

Table (14): Duncan statistical test for the four weeks control subgroup

	Mean	Standard deviation	Duncan
0WP	0.7833	0.1472	A
2WP	0.9667	0.05164	B
4WP	0.8833	0.07528	AB

DISCUSSION

No significant literature exists comparing the effect of systemic bisphosphonate therapy on endosseous implant osseointegration. Some studies have concentrated on locally applying bisphosphonates to implant surfaces and then measuring the bone response. Statistically significant increases in bone density and bone formation occurred with the alendronate-coated implants.⁽¹⁷⁾

This study showed that in general no significant differences between the treated and plane subgroups according to the treatment course given in that the stability measurements of the treated subgroups before loading of the micro-screw implants were different, although not significant which larger in 0WT subgroup. This probably due to the effect of primary stability, then 4WT subgroup may represent the beginning of increase bone density around the micro-screw implant and the lower stability of 2WT subgroup could be due to the process of bone remodeling (resorption and apposition) which ordinarily occur as a healing process may affect its stability, unlike the control group which showed a significant difference. Same difference after loading but significant for 0WT subgroup and not for 2WT and 4WT subgroups. For 0WT subgroup, it is probably the result of increase fitness (a result of added tension of loading) and add to the influence of primary stability, but for 2WT subgroup as described above it may be due to the remodeling process and for 4WT subgroup, here the effect of loading may do breaking of initial formed bone attachment, this result same that of control group.

The stability measurement of the three treated subgroups after a healing periods of two weeks for the loaded micro-screw implant showed a differences, but not significant which higher for 0WT subgroup which could be due to the effect of primary stability add to spring tension from zero

to two weeks period then 4WT subgroup, which probably be due to increase bone formation around the screw due to healing period of 6 weeks that increase its fitness, this come in accordance with the result of control subgroup.

The stability measurement for 0WT subgroup showed no significant differences from time of instillation of screw till loading then 2 weeks after loading. This could be the result of a short time and still under the control of primary stability. This is not in accordance with the control subgroups. For 2WT subgroup, a significant difference had been shown from time of instillation, 2 week then loading, 2 week after loading, with stability higher for the first time of instillation and this is already explained, same result for control subgroups. similar to 4WT subgroups a significant difference in mean from time of instillation, 4 week then loading, 2 week healing period after loading, although the stability mean for 4WT subgroup better than 2WT subgroup which may be due to the time factor. The screw may need time after losing the primary stability and getting the secondary one by bone cell accumulation around the screw (increase bone density or possibly osseointegration), this is also similar to those control subgroups.

In general, we notice that the initial reading of stability is higher than subsequent measurement although time factor is important that is to say with the time and progress of bone healing around the screw will probably lead to bone cell accumulation around the screw, but here the size of screw may affect the stability, because we use the smaller sort in the micro-screw implant kit. This mean that we may need a longer time for the smaller diameter to establish a good secondary stability and comparing to other studies on micro-screw all using nearly double. This size also may get no benefit of using one course of multiple course recommend by the manufacture that subjecting the patient to a multi-

ple courses of medication is a legal point of view which even in one course and also need time which is not problem in dental implant, but for micro-screw implant is a time temporary process of nearly maximum one year so we may finish the use of the implant and still need time to finish the course of medication.

Saito ⁽¹⁸⁾ suggested that orthodontic force should be loaded on the micro-screw after 18 weeks of healing. Roberts ⁽¹⁹⁾ concluded that micro-screws could stand orthodontic loading of 100 g after 6 weeks of healing. Studies by Costa ⁽²⁰⁾ indicated that micro-screws could provide stable anchorage after 4 weeks of healing. Melsen ⁽²¹⁾ reported that osseointegration could be observed on the immediately loaded bone-implant interface. Some clinicians suggest that some healing time is required and recommend delaying force application. ⁽²²⁾ Others, however, state that orthodontic force can be applied immediately after implantation. ⁽²³⁾

A complete understanding of this drug class and the effects on long-term implant osseointegration in humans will require further study.

CONCLUSION

According to the result of this study all the stability in different time interval are acceptable according to the stability standard criteria of the periotest prescribed before, thus micro-screw implant even smaller diameter can be used as a fixed anchorage in orthodontics and possibly can be loaded safely from immediate instillation.

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