MACRO VERSUS MICRO THREAD DESIGN IMPLANT AND THEIR EFFECT ON THE STABILITY

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ABSTRACT

INTRODUCTION: Dental implants are considered stable tools for replacing missing teeth, achieving long-term success rates above 90 percent, and their use in dental practice has become normal. One of the reasons to this success is the primary reliability of an implant. Implant design is one of the main factors that play a role in stability.

OBJECTIVES: The purpose of this study was to compare the role of the Macro thread design and the Micro thread design on implant stability through the use of the Resonance Frequency Analysis (RFA).

MATERIALS AND METHODS: The study was carried out on 6 patients, each of these patients has missing teeth on each side of posterior area in the lower jaw. All the right sides received Micro thread design implant (dentium implant,) and the left side received Macro thread design implant (Megagen AnyRidge). After dental implants were placed in their sites, the stability was measured by using the Resonance Frequency Analysis (RFA) to assess the stability of the two types of implants at three times periods: At the time of placing the implant, 3 months and 6 months.

RESULTS: The mean implant Stability value for Group A was 70.57±5.74 immediately post-operatively, on the 3 month to 77.14±6.74 and reach 84.29±6.02 on the 6 month, for Group B was 63.29±5.85 immediately post-operatively, on the 3 month 70.57 ± 4.69 and on the 6 month 77.14±4.53. The mean bone density values for Group A at 3 months was 481.98± 51.78 and at 6th month was 504.28± 47.50, in Group B the 3 months was 439.54±70.49 and at 6th month was 463.83± 74.44.

CONCLUSION: The Micro thread design implants shows higher stability than Macro thread design.

KEYWORDS: Osseo integration, Implant design, Dentium implant, Megagen AnyRidge, Resonance Frequency designs available Analysis, CBCT.

RUNNING TITLE: Macro, micro thread design, implant stability.

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INTRODUCTION

Dental implants are considered one of the most well-known effective ways of missing teeth replacement. Realizing success rates more than 90% in the long follow-up period, and their use has become familiar in dental practice. The first step of successful osseointegration is the primary stability of an implant (1).

In the 1960s the modern dental implant has become one of the critical choices of the treatment option in the replacement of missing natural teeth. Since then, the dental implant industry has recently seen enormous development in a vast number of manufacturers and several designs are available. Recently, a variety of implant lengths, surfaces, body designs, platform connections, thread forms, and body designs are available, where these variations in implant designs available can aid in primary stability (initial stability) and Osseointegration. It has been reported that implant design is a vital parameter for obtaining primary stability (2). Also, the presence of Osseointegration is crucial to evaluate implant success (3).

Various mechanical factors form the effective connection between an implant and its surrounding bone. One of these factors is the design of implants which Specifies stability and distribution of pressure during the process of osseointegration (3), where the implant designs contain macroscopic features include (body design and thread geometry) and the microscopic features include (implant materials, surface morphology, and surface coatings).

The optimal design of the implant itself can increase the efficiency of the Osseointegration process. The implant stability as well as the structural features of an implant effect on primary contact to improve the implant's initial stability. It plays a major role in developing the ability to resist forces during the Osseointegration process (3).

There are many factors have not been adequately studied that could have effects on bone remodeling in the postoperative period, and implant thread design is one of these factors, including thread pitch, depth, and shape e.g., V-shape, reverse buttress, block) (4).

Several studies have shown various surface features that play an essential role in the implant during the initial stage of bone integration. Macro irregularities such as grooves and pores of different dimensions, known to be the parameters of implant design. Implant threads were introduced to improve the initial contact with the bone, increase the surface area and thus dissipate interfacial tension (5). Other studies have also stated the thread designs resulting in superior bone response to that surface with modification of the implant surface treatment methods (5).

Aim of the study is comparing between the role of Macro thread design and the Micro thread design on implant stability through use Resonance Frequency Analysis (RFA).
This study proposes that this Macro thread design may have similar results to Micro thread design in their effect on the stability of the implant.

MATERIALS AND METHODS
Appropriate ethical clearance was obtained from the institution at which the study was conducted. Where all patients received thorough explanations about the planned treatment and its potential risks and complications and signed a written informed consent form before being enrolled in the study. It was also mentioned that the patient had the right to withdrawal from the study anytime without any consequences. Ethical approval for this study was obtained from the research ethics committee, Faculty of Dentistry, Alexandria University before beginning the study.

Study design
The present study was a randomised controlled clinical trial (split mouth design) performed in six patients with 14 dental implants. Each patient in attendance received one implant on each side, except for one of those patients who received two implants on each side. (Seven of the implants were Macro thread design Megagen Any Ridge and the other seven were Micro thread Dentium implants). The age range of the participating patients was between 30-50 years of age. All cases of missing mandibular teeth indicated for implant placement by two-stage surgery. Patients have been chosen from the Outpatient Clinic of the Department of Oral and Maxillofacial Surgery, Faculty of Dentistry, University of Alexandria.

The dental implants were divided into two groups (A and B):
- **Group (A):** On the right side consist of 7 implants with Micro thread design.
- **Group (B):** On the left side consist of 7 implants with Macro thread design.

Criteria for patient's selection
- **Inclusion criteria:** A healthy condition that initiates bone healing, a mandibular edentulous site for implant placement and a sufficient volume of bone at the implant placement site.
- **Exclusion Criteria:** Systemic illness, drug or habit considered to have a negative effect on bone healing and/or dental implant success: (poorly controlled diabetes [HbA1c > 8%], heavy smoker, para dental habits, history of use of bisphosphonate, history of radiation therapy of the head and neck affecting the proposed implant site, current use of medication adverse to healing (e.g. corticosteroids, chemotherapy).

Materials
- **Group (A):** Dentium Implant System (Super Line)
  - Micro thread design was placed on the right side.
  - Tapered design, Extended cutting edge, Double thread design.
  - The size of the implant that was placed in this study 4.0 mm diameter and 10mm length.
- **Group (B):** Megan Implant System (AnyRidge)
  - Macro thread design was placed on the left side.
  - Incorporates a novel thread design, which includes rounded, non-cutting edge, wide thread depth, and increased thread pitch compared to a conventional thread design.
  - The size of the implant that was placed in this study 4.0 mm diameter and 10mm length.

Surgical kit
This includes the surgical instruments used for flap reflection, an electric motor with irrigation system, drills with different sizes for each system, guide drill, drill extension countersink, depth gauge, and parallel pins, drivers, hand wrench, ratchet wrench, a contra-angle handpiece, and Osstell.

Pre-operative Stage
Each patient was evaluated by taking appropriate history and thorough clinical examination. Initial periodontal treatment was completed, including scaling and oral hygiene training.

Preliminary assessment
For both arches and diagnostic test designs, the primary alginate impression was taken, clinically as well as on the study design, the inter-arch relationship, interocclusal space that could accommodate the implant abutment, and potential crown restoration were evaluated. Vacuum-Formed surgical guidance stent was manufactured and in the stent opposite the missing tooth a hole was drilled indicating the implant's location.

**Radio-graphical examination**
- For the examination of bone height, panoramic radiography was obtained to detect the presence of any pathology at the implant site.
- Cone-beam computed tomography (CBCT) was done to detect (buccolingual width of the ridge, length, and diameter of the implant, bone density, approximation to vital anatomical structures. (Figure 1)

**Surgical Procedures**
The two groups were operated under local anesthesia (mandibular nerve block technique), using Articaine with vasoconstrictors (1:200 000).
- Mid crestal incision was done down to the bone with a blade No.15 was utilized, including one tooth on either side of the proposed implant site. (Figure 2A, 3A)
- Sharp dissection was performed by a perioisteal elevator.
- Implant placement procedures are done in compliance with the manufacturer's instructions.
- The electric motor was used to prepare the implant site together with a high-speed high torque handpiece externally irrigated.
- Sterile saline was used during the preparation of the implant site for external irrigation.
- The preparation of the bone site was undertaken using the 1 mm pilot drill with external irrigation. It adjusted the electric motor to 1000 rpm.
- The initial drill 2 mm was used to reach to the predetermined length.
- Sequential drilling has been done up to the final drill. (Figure 2B, 3B)
- The depth gauge and path indicator were used to assess the osteotomy site depth and to test the implant's parallelism. (Figure 2C, 3C)
- The implant was inserted manually into its bed under moderate apical pressure until it stopped. Then the implant was inserted into the final mounting depth by a ratchet wrench. (Figure 2D, 3D)
- Osstell smart peg connected to the fixture and initial stability measurement taken. (Figure 2E, 3E)
- Cover screw was inserted and secured into the implant's occlusal opening. (Figure 2F, 3F)
- Flap repositioned after the surgical site was irrigated and dry.
- 3.0 black silk suture material used to suture the flap.

**Figure (1):** (a) Pre-operative CBCT- (b) Radiograph after 3 months postoperative CBCT- (c) Radiograph with final restoration after 6 months postoperative.

**Figure (2):** (a) Mid Crestal Flap Reflection. (b) Drilling the bone. (c) Parallel Pin. (d) Implant placement (Dentium dental implant) Group A. (e) Osstell measuring. (f) Implant with cover screw. (g) Abutment in place right side (Dentium dental implant). (h) Final porcelain restoration lower right six.

**Figure (3):** (a) Mid Crestal Flap Reflection. (b) Drilling the bone. (c) Parallel Pin. (d) Implant placement (Maga-Gan dental implant) Group A. (e) Osstell measuring. (f) Implant with cover screw. (g) Abutment in place right side (Maga-Gan dental implant). (h) Final porcelain restoration lower left six.

**Post-surgical phase**

The patient was instructed to rinse their mouth with antiseptic chlorhexidine mouthwash 3 times a day starting from the second day postoperatively and continued for 2 successive weeks.

Postoperative panoramic x-rays were taken to ensure proper placement of the implant and the implant's relationship to opposing landmarks or surrounding structures.

**Follow up phase**

The following clinical parameters were recorded immediately after loading (baseline) and at the end of the study:

1. **Presence or absence of pain or sensitivity**

   The patient comfort was assessed using the Visual Analogue Scale (VAS) (6) of 0 to 10.

2. **Probing depth (PD)**

   According to Harvard Conference (7), the probing pocket depth around the implant was measured on the four dimensions of the implant's facial, palatal and proximal surfaces using a mm periodontal probe graduation.

3. **Healing index score**

   HI recording was done on the post-surgery first-week. The HI rates healing based on redness, tissue appearance, swelling, suppuration, and epithelialization.

   Landry, Turnbull, and Howley (8) have developed a healing index to explain the degree of medical healing following periodontal surgery, and it has also been recently updated to be used for socket healing. For post-extraction score (0/1) with a total score of seven, with a total score of 7: presence/absence of redness; presence/absence of granulation tissue; presence/absence
of suppurition; presence/absence of swelling; degree of tissue epithelialization (partial/complete); presence/absence bleeding; presence/absence of pain on palpation.

IV. Stability Evaluation
When evaluating implant stability and osseointegration, the Osstell was measuring stability at three and six months. The result was provided as a 1-100 Implant Stability Quotient (ISQ) value (9,10). Osstell is a non-invasive procedure first used in 1996. This consists of a small L-shaped transducer fastened to the implant or the transmucosal abutment using a screw. There is a vertical beam connected to this transducer with two piezoceramic components. One of the piezoceramic elements produces a vibration consisting of a small, 5 to 15 kHz sinusoidal signal in 25 Hz steps. The other piezoceramic component analyzes the transducer's response to the vibration. The higher the ISQ the more stable is the implant.

V. Radiographic Evaluation
Cone Beam Computed Tomography (CBCT): is a three-dimensional imaging technology that offers less radiation dosage reportedly up to 15 times lower than those of conventional CT scans.
Assessment of bone density after three and six months from dental implants placement using CBCT, after dental implant placement bone density was measured around the dental implant site by using Hounsfield unit HU to detect the osseointegration. Three controlled and standardized dimension square areas were selected just mesial, distal and apical to the implant including the Bone-implant interface, Mean, Standard deviation, Minimum and Maximum readings were automatically displayed by the system "On-demand 3D" and used for statistical analysis.

Prosthetic phase
The cover screw was removed and the healing abutment was tightened at the third month. (Figure 2G, 3G)

After 4 months of this study the patients were recalled for delivery of the definitive restoration, which impressions were taken and sent to the laboratory for the fabrication of the final restoration. (Figure 2H, 3H)

Statistical analysis
Statistical analysis Data were fed to the computer and analyzed using version 20.0 (11) of the IBM SPSS software package (Armonk, NY: IBM Corp). Using range, mean, standard deviation and median, quantitative data were described. By the Kolmogorov-Smirnov method, the distribution of quantitative variables was checked for normality. The paired t-test was used to compare two periods of quantitative variables normally distributed, while the ANOVA of repeated measurements was performed. Collecting and tabulating data. The mean Probing depth scores of the control group were 1.64 ± 0.56 with a maximum recorded value of 2.0, while the mean Probing depth scores of Group A were 1.36 ± 0.38 with a minimum recorded value of 1.0 and a maximum recorded value of 2.5. This difference in the probing depth score (P2~<0.001) was found to be statistically significant. The mean probing depth scores of Group A on the sixth month were 1.36±0.38 with a minimum reported value of 1.0 and a maximum recorded value of 2.0, while the mean probing depth scores of the control group were 1.64 ± 0.56 with a minimum recorded value of 1.0 and a maximum recorded value of 2.5. This difference in the probing depth score (P2~<0.001) was found to be statistically significant. (Figure 4 A, B) (Table 1)

Table (1): Comparison between the studied groups according to peri-implant probing depth throughout the study period.

<table>
<thead>
<tr>
<th>Peri-implant probing depth</th>
<th>Group A (n = 7)</th>
<th>Group B (n = 7)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>3rd month</td>
<td>6th month</td>
</tr>
<tr>
<td>Min. – Max.</td>
<td>1.0 – 2.50</td>
<td>1.0 – 2.0</td>
</tr>
<tr>
<td>Mean ± SD.</td>
<td>1.82 ± 0.47</td>
<td>1.36 ± 0.38</td>
</tr>
<tr>
<td>Median</td>
<td>2.00</td>
<td>1.50</td>
</tr>
</tbody>
</table>

III. Healing index score
Healing index score was evaluated from the first week till a month, none of the implants showed any (redness, granulation tissue, suppurition, swelling and bleeding) score was (0).

IV. Implant Stability Evaluation
At the time of implant insertion, the implant stability measurement was examined 3 times, 3rd month and 6th month for both groups using the Resonance Frequency Analysis through the Osstell (ISQ) system.

Group (A):
The implant stability for patients of (Group A) who received Micro thread implant placement and the results were measured by using the Resonance Frequency Analysis.

Their ages ranged from 30 to 50 years, the teeth that this study replaced were multi-rooted mandibular teeth, all patients were free of any local or systemic infection.

The 14 implants were divided into two groups, 7 implant Macro thread design: (Mega Gen) and 7 implants Micro thread design (Dentium)

Group (A): Included seven Micro thread implants that were placed on the right side of the lower jaw.

Group (B): Included seven Macro thread implants that were placed on the left side of the lower jaw.

Clinical evaluation
I. Presence of pain
Using visual analog scale (VAS) from 0 to 10 ("0" is pain-free and ”10” is unsustainable pain), the pain was evaluated daily for first week. After surgery, patients reported mild pain at the surgical site, which had gradually subsided after the procedure by the fourth day.

II. Probing depth
Probing depth was measured for all implant axial surfaces, statistical analysis of sampling depth scores for all patients was performed. Collecting and tabulating data. The mean Probing depth scores of group A on the third month were 1.82 ± 0.47 with a minimum documented value of 1.0 and a maximum recorded value of 2.5, while the mean Probing depth scores of group B were 2.00 ± 0.65 with a minimum recorded value of 1.50 and a maximum recorded value of 2.50. This difference in the probing depth score (P2~<0.001) was found to be statistically significant. (Table 4 A, B) (Table 1)
From the table above, it's shown that the maximum stability measured at the time of operation was of (ISQ) = 80, (ISQ) units and the minimum measured was 65 (ISQ) units with an average mean and standard deviation = 70.57 ± 5.74.

Then after 3 months, it was shown that the maximum stability measured was of (ISQ) = 74 (ISQ) units and the minimum measured was 68 (ISQ) units with an average mean and standard deviation = 77.14 ± 4.69.

After 6 months, the maximum stability measured was of (ISQ) = 77 (ISQ) units and the minimum measured was 71 (ISQ) units with an average mean and standard deviation = 70.57 ± 4.69.

After 6 months, the maximum stability measured was of (ISQ) = 77 (ISQ) units and the minimum measured was 65 (ISQ) units with an average mean and standard deviation = 70.57 ± 5.74.

Comparing the above stability results for the two groups (A & B), it was shown that the average mean and standard deviation of the stability for all Group A cases receiving Macro thread design implants were 77.33 ± 5.93 ISQ units. That is greater than the average mean and standard deviation = 70.57 ± 4.69.

After 6 months, the maximum stability measured was of (ISQ) = 78 (ISQ) units and the minimum measured was 74 (ISQ) units with an average mean and standard deviation = 74.44.

Comparing the above-stated bone density findings for the two groups analyzed, it was shown that the average mean and standard deviations in bone density for all Group A cases receiving Macro thread design implants were 77.33 ± 5.93 ISQ units. That is greater than the average mean and standard deviation = 70.57 ± 4.69.

After 6 months, the maximum bone density measured was 572.83 and the minimum measured was 370.57 with an average mean and standard deviation = 463.83 ± 74.44.

### Table (2): Comparison between the two studied groups according to stability.

<table>
<thead>
<tr>
<th>Stability</th>
<th>Group A (n = 7)</th>
<th>Group B (n = 7)</th>
<th>t</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>1&lt;sup&gt;st&lt;/sup&gt; Month</td>
<td>Mean ± SD.</td>
<td>70.57 ± 5.74</td>
<td>63.29 ± 6.58</td>
<td>3.43</td>
</tr>
<tr>
<td>Median</td>
<td>70</td>
<td>62</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Min – Max</td>
<td>65-80</td>
<td>56-74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2&lt;sup&gt;nd&lt;/sup&gt; Month</td>
<td>Mean ± SD.</td>
<td>77.34 ± 6.74</td>
<td>70.57 ± 4.69</td>
<td>2.35</td>
</tr>
<tr>
<td>Median</td>
<td>79</td>
<td>71</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Min – Max</td>
<td>68-84</td>
<td>65-77</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3&lt;sup&gt;rd&lt;/sup&gt; Month</td>
<td>Mean ± SD.</td>
<td>84.29 ± 6.02</td>
<td>77.14 ± 4.53</td>
<td>3.28</td>
</tr>
<tr>
<td>Median</td>
<td>86</td>
<td>78</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Min – Max</td>
<td>69-85</td>
<td>65-77</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean ± SD.</td>
<td>77.33 ± 5.93</td>
<td>70.33 ± 4.29</td>
<td>3.999</td>
<td>0.007</td>
</tr>
<tr>
<td>Median</td>
<td>78.3</td>
<td>69.33</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average</td>
<td>70.57 ± 5.74</td>
<td>70.57 ± 5.74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean ± SD.</td>
<td>74.44</td>
<td>74.44</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>78</td>
<td>69</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Radiographic Evaluation

#### Bone density

The bone density was measured using the On demand program. The mean was calculated after 3, 6 months; show that there was a significant increase in the mean bone density by time at the two periods of follow up in both groups for all cases.

**In group A**

The maximum bone density measured after 3 months was 532.8 and the minimum measured 388.43 measured with an average mean and standard deviation = 481.98 ± 51.78.

After 6 months, the maximum bone density measured was 582.8 and the minimum measured was 442.47 with an average mean and standard deviation = 504.28 ± 47.50.

**In group B**

The maximum bone density measured after 3 months was 561.53 and the minimum measured 384.03 measured with an average mean and standard deviation = 439.54 ± 70.4.

After 6 months that the maximum bone density measured was 572.83 and the minimum measured was 370.57 with an average mean and standard deviation = 463.83 ± 74.44.

The average value of all cases of group A, mean bone density value was 451.69, with SD 70.09. Cases of group B mean bone density value was increased to 493.13, with SD 47.35.

### Table (3): Mean bone density values for different periods for group (A&B).

<table>
<thead>
<tr>
<th>Bone density</th>
<th>Group A (n = 7)</th>
<th>Group B (n = 7)</th>
</tr>
</thead>
<tbody>
<tr>
<td>3&lt;sup&gt;rd&lt;/sup&gt;</td>
<td>481.98 ± 47.50</td>
<td>439.54 ± 47.50</td>
</tr>
<tr>
<td>6&lt;sup&gt;th&lt;/sup&gt;</td>
<td>451.69 ± 51.78</td>
<td>493.13 ± 47.35</td>
</tr>
<tr>
<td>Mean ± SD.</td>
<td>388.43 ± 504.28</td>
<td>384.03 ± 439.54</td>
</tr>
<tr>
<td>Median</td>
<td>532.8</td>
<td>561.5</td>
</tr>
<tr>
<td>Min – Max</td>
<td>384.03</td>
<td>493.13</td>
</tr>
<tr>
<td>Average</td>
<td>481.98</td>
<td>493.13</td>
</tr>
</tbody>
</table>
DISCUSSION
Stability is one of the key factors in the successful treatment of dental implants. Branemark's osseointegration, meaning the structural and functional coordination between the implant and the underlying vital bone, is focused on long-term endosseous dental implant.

Bone density and implant stability are important factors for implant Osseointegration which have been widely demonstrated by several authors. It is suspected that the lack of primary stability is the cause of early implant failure (12). Stability depends on bone strength, surgical procedure and the micro- and macro-design of the implant used; thus, implant stability is the key to clinical success (13-15).

The present study aimed to compare two different implant profiles Micro thread "Narrow pitch" implants and Micro thread "wide pitch" implant was tested in the posterior area of the lower jaw, where we use this study to evaluate the influence of implant thread design on stability and osseointegration process. The purpose of this hypothesis is that selecting implant design features that increase surface area for contact that may improve the stability and bone-implant contact.

In this study, six patients with missing mandibular posterior teeth were chosen from the Outpatient Clinic, Faculty of Dentistry, Alexandria University, Department of Oral and Maxillofacial Surgery. The chosen patients were free from any untreated systemic disorders or conditions that might hinder the surgical or healing process of the implant.

Bornstein, Cionca et al., (16) investigating whether systemic diseases with / without systemic medicines increased the risk of implant failure and thus decreased the effectiveness and survival of dental implants, stated that the level of evidence indicating absolute and relative implant therapy contraindications due to systemic diseases was small.

Patients with bruxism, heavy smokers, chemotherapy or radiotherapy patients, and patients with immune suppression were the ones omitted from this study. Gómez-de Diego et al. (17) conducted this study to analyze clinically affected patients' indications and contraindications of dental implants and concluded that the use of nicotine and head and neck radiotherapy is associated with increased loss of dental implants.

CBCT was used to measure bone density due to its sensitivity, lower radiation exposure and lower cost compared to CT.

The mean bone density in our sample increased significantly towards the post-operative six-month period. The results show that there was a significant increase in the mean bone density by time at the two periods of follow up in both groups for all cases:

The average value of all cases of group A, mean bone density value was 493.13, with SD 47.35, the Average value of all cases of group B, mean bone density value was increased to 451.69, with SD 70.09.

However, in 2011, Naser et al. (18) compared continuous films taken over time at different time scales to research alveolar bone changes and bone density around dental implants, finding that the average density obtained at different standard densitometry stages showed a gradual increase in bone density throughout the phase.

Implant stability was tested through the Osstell ISQ method using Resonance Frequency Analysis (RFA), RFA was chosen as a non-invasive and reliable approach to assess variation over time in implant stability. RFA registrations are directly related to the stability of the implant in the surrounding bone: new bone apposition at the implant-bone interface may be expressed during healing and increase of the ISQ values (19-20).

Figure (4): (a) Evaluation of probing depth for the 3 Months period for Group (A) and (B), (b) Evaluation of probing depth for the 6 Months period for Group (A) and (B).

Figure (5): Comparison between mean stability values for different periods for groups (A&B).

Figure (6): Comparing between the bone density values for different periods for group (A&B).
The normal range of ISQ values commonly recorded for primary stability implants is between 60 and 80, although a consensus has not been developed on the ISQ threshold below which an implant should not be considered stable. Meredith et al. (21) concluded that RFA is a method that can serve as a valuable research technique and is useful in studying the operation of implants in the surrounding tissue. Jaramillo et al. (22) have stated that in Osstell Mentor and Osstell ISQ, Resonance frequency analysis systems show almost perfect reproducibility, repeatability, and accuracy. Results from this study showed that "Narrow pitch" Micro thread implants obtained higher primary stability values compared to "wide pitch" Macro thread implants measured with RFA. Our result shown the average value of stability measured for all group A cases, Min.-Max = 69-85 (ISQ) units, the overall mean and standard deviation are 77.33 ± 5.93 (ISQ) units for all group A cases and the average value of all cases of group B, Min. – Max =65-77 (ISQ) units, the average mean and standard were 70.33±4.29. These results were in accordance with similar studies which discussed similar relation and found that smaller pitch implant design increased bone-implant contact and primary stability. Also, similar results were obtained by other studies. Implants with smaller pitch are beneficial by increasing BIC (23).

In 2012, Lan et al., (24) found that the loading form is the main factor affecting the pressure distribution and that the thread pitch above 0.8 mm is more appropriate for screwed implants in biomechanical consideration. Kong et al., (25) thread pitch plays a greater role than buccolingual load in protecting dental implants under axial load. The optimum configuration in a cylinder implant should be the thread pitch exceeding 0.8 mm, but excessive pitch should also be avoided. Abuhussein et al., (26) conclude that attaching threads or micro threads to an implant's crystal module may potentially increase bone-implant contact and primary stability. Also, similar results were obtained by other studies. Implants with smaller pitch are beneficial by increasing BIC (23).

CONCLUSION

According to this study, the geometry of dental implant threads affects the stability and BIC of the Micro thread design implants shows higher stability (as calculated using ISQ) than the Macro thread design. On the other hand, the bone density of the Micro thread design appears slightly higher than that of the Macro thread design.

CONFLICT OF INTEREST

The authors declare that they do not have any conflicts of interest.

REFERENCES