

Original Research

# Influence of implant design, length, diameter, and anatomic region on implant stability: a randomized clinical trial



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## ABSTRACT

**Objectives:** To evaluate the influence of implant geometry and anatomical region on implant stability.

**Methods:** A randomized controlled clinical trial was conducted on 45 patients, in whom a total of 79 implants were placed: 40 MIS C1 Implants and 39 MIS Seven Implants. The implant stability quotient was measured using resonance frequency analysis immediately after implant placement and 8 weeks later with an Osstell Mentor device.

**Results:** 76 implants were analyzed. The implant stability quotient was statistically significantly higher for secondary stability than primary stability ( $68.7 \pm 8.6$  vs.  $65.2 \pm 10.3$ , respectively,  $p=0.023$ ). Considering primary stability, no statistical differences were found between the implant lengths 8.0 mm, 10.0 mm, 11.0 mm, and 11.5 mm ( $67.9 \pm 7.6$ ,  $63.9 \pm 10$ ,  $57.2 \pm 11.1$ , and  $66.4 \pm 11.3$ , respectively,  $p=0.312$ ). The same was observed for secondary stability ( $68.4 \pm 9.4$ ,  $67.9 \pm 9.3$ ,  $74.7 \pm 1.5$ , and  $69.2 \pm 7.9$ , respectively,  $p=0.504$ ). Also, there were no statistically significant differences between the implant diameters 3.75 mm and 4.20 mm concerning primary stability ( $64.3 \pm 8.7$  and  $66.1 \pm 11.7$  respectively,  $p=0.445$ ) or secondary stability ( $68.8 \pm 8.2$  and  $68.7 \pm 9.1$  respectively,  $p=0.930$ ). Regarding implant design, a statistically significant difference was found only for secondary stability, favoring MIS Seven implants ( $p=0.048$ ). The intraoral location was statistically significant for both primary and secondary stability, as these were higher on the anterior maxilla than the posterior maxilla and mandible ( $p<0.05$ ).

**Conclusions:** The diameter and length of the implants studied did not influence their stability. Implant design may influence secondary stability, whereas intraoral location has a relevant effect on primary and secondary stability. (Rev Port Estomatol Med Dent Cir Maxilofac. 2021;62(1):xxx-xxx)

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## Influência do comprimento, diâmetro e desenho do implante e da região anatômica na estabilidade implantar: ensaio clínico randomizado

### R E S U M O

#### Palavras-chave:

Desenho do implante  
Diâmetro do implante  
Comprimento do implante  
Estabilidade do implante  
Osseointegração  
Análise de frequência de ressonância

**Objetivos:** Avaliar a influência da geometria dos implantes e região anatômica na estabilidade implantar.

**Métodos:** Foi realizado um ensaio clínico controlado aleatorizado em 45 pacientes, onde foram colocados 79 implantes: 40 implantes MIS C1 e 39 implantes MIS Seven. O coeficiente de estabilidade do implante foi medido através da análise de frequência de ressonância (RFA) imediatamente após a colocação do implante e 8 semanas depois, com Osstell Mentor.

**Resultados:** Foram analisados 76 implantes. O coeficiente de estabilidade do implante registrado para estabilidade secundária foi significativamente maior que o verificado para a estabilidade primária ( $68,7 \pm 8,6$  e  $65,2 \pm 10,3$ ,  $p=0,023$ ). Considerando a estabilidade primária, não foi encontrada diferença estatisticamente significativa entre os comprimentos de 8,0 mm, 10,0 mm, 11,0 mm e 11,5 mm ( $67,9 \pm 7,6$ ,  $63,9 \pm 10$ ,  $57,2 \pm 11,1$  e  $66,4 \pm 11,3$ ,  $p=0,312$ ). O mesmo foi verificado para a estabilidade secundária ( $68,4 \pm 9,4$ ,  $67,9 \pm 9,3$ ,  $74,7 \pm 1,5$  e  $69,2 \pm 7,9$ ,  $p=0,504$ ). Não se observou diferença estatisticamente significativa entre os diâmetros dos implantes de 3,75 e 4,20 mm e estabilidade primária ( $64,3 \pm 8,7$  e  $66,1 \pm 11,7$ ,  $p=0,445$ ) ou estabilidade secundária ( $68,8 \pm 8,2$  e  $68,7 \pm 9,1$ ,  $p=0,930$ ). Foi encontrada diferença estatisticamente significativa para a estabilidade secundária, favorecendo o implante MIS Seven ( $p=0,048$ ). A localização intraoral foi estatisticamente significativa para a estabilidade primária e secundária, sendo maior na maxila anterior em relação à maxila e mandíbula posteriores ( $p<0,05$ ).

**Conclusões:** O diâmetro e o comprimento dos implantes estudados não influenciaram sua estabilidade. O tipo do implante parece influenciar a estabilidade secundária, enquanto a região anatômica aparenta ter um efeito relevante na estabilidade primária e secundária. (Rev Port Estomatol Med Dent Cir Maxilofac. 2021;62(1):xxx-xxx)

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### Introduction

The use of dental implants is a widespread treatment option for the replacement of missing teeth.<sup>1</sup> This treatment's predictability is directly related to osseointegration, which has been defined as the direct structural and functional connection between bone and the surface of a dental implant.<sup>2</sup> Some of the most relevant factors for the osseointegration of dental implants are achieving and maintaining implant stability.<sup>2</sup> Insufficient implant stability has been cited as the greatest potential risk factor for implant loss.<sup>3,4</sup>

The implant stability can be measured in two different stages: primary and secondary stability.<sup>5</sup> Primary stability is the initial biometric stability obtained immediately after implant insertion that results from the mechanical attachment within the bone and seems to be affected by bone quality and quantity, surgical technique, and implant geometry.<sup>3</sup> Secondary stability offers biological stability through bone regeneration and remodeling and seems to be highly affected by primary stability.<sup>6</sup>

The objective measurement of implant stability is important to evaluate the clinical outcome of dental implant treatments throughout implant healing and loading. There are several methods to assess implant stability that can be grouped as invasive and non-invasive.<sup>3</sup> The invasive methods include

histologic and histomorphologic analysis, tensional test, push- and pull-out test, and removal torque analysis.<sup>3,7</sup> The non-invasive methods for assessing implant stability include radiographic analysis with imaging techniques, cutting torque resistance, insertion torque measurement, reverse torque, modal analysis, percussion test, pulsed oscillation waveform, resonance frequency analysis, and magnetic technology.<sup>3</sup>

Developed by Meredith et al. in 1996, the resonance frequency analysis (RFA) is a reliable and easily predictable non-invasive objective method for quantifying primary and secondary implant stability.<sup>8,9</sup> It measures implant stability as a reaction to oscillations exerted onto the implant/bone interface, providing a unit of measurement known as the implant stability quotient (ISQ), expressed in a range from 0 to 100.<sup>5</sup> According to several studies, RFA is a highly effective non-invasive method to determine dental implants' stability during the healing and follow-up treatment stages.<sup>10-12</sup> Previous research reported that ISQ levels for successfully osseointegrated implants fluctuate between 57 and 82, whereas implants with ISQ values lower than 50 have a high risk of failure.<sup>13</sup>

Published results on the relationship between implant stability, design, length, and diameter are controversial. Previous research has shown that lower stability was obtained with longer implant lengths.<sup>14</sup> Other investigations obtained dif-

ferent results, concluding that implant length has no significant effect on implant stability.<sup>15</sup> However, those authors concluded that ISQ values increased with increased implant diameter, and tapered implants showed higher ISQ values than parallel implants. Given the diversity of the current results, it is important to carry out a clinical trial that simultaneously studies the different factors that appear to influence implant stability.

Therefore, this randomized controlled clinical trial aims to evaluate the influence of implant length, diameter, design, and anatomic region on ISQ values during osseointegration. The research hypothesis was that ISQ values are affected by jaw region, implant diameter, implant length, and implant design.

## Material and Methods

A prospective, randomized, parallel-group, controlled clinical trial was carried out at the Implant Dentistry Postgraduate Program of the Faculty of Dental Medicine of the University of Lisbon (FMDUL), Lisbon, Portugal. Inclusion criteria were adults with good general health, non-smokers, with at least one tooth missing for more than 6 months and sufficient bone height/width for implant placement without the need for bone augmentation. Exclusion criteria included patients with uncontrolled systemic diseases, history of head/neck irradiation, immune system severe deficiencies, smokers, and patients treated with oral/intravenous bisphosphonates.

Using G\*Power (version 3.1.9.4, Heinrich-Heine-University, Germany), a minimum sample size of 70 implants was calculated as required to detect comparisons between the two groups, considering a statistical power of 80% and estimated dropouts. The eligible patients were randomly allocated into two different groups by simple randomization (using GraphPad Software, QuickCalcs – website <http://www.graphpad.com>) according to implant design: Group A – C1 implants (MIS Implants Technologies Ltd) and Group B – Seven implants (MIS Implants Technologies Ltd).

The study was approved by the Ethics Committee of the FMDUL, Lisbon, Portugal. All procedures followed the standards of the Helsinki Declaration of 1975, as revised in 2000. All patients were fully informed of the nature of the study and signed an informed consent form.

A total of 79 implants were placed: 40 MIS C1 implants and 39 MIS Seven implants (MIS Implants Technologies Ltd). All implants had the same surface: sandblasted and acid-etched surface. C1 implants feature a 12-degree conical connection that ensures a secure fit between abutment and implant. The C1 implant has a dual thread design (range 1.5 mm) over the entire implant body. The apical region of C1 implant features two cutting blades establishing the self-tapping properties. Seven implants' apical region features an internal hexagon connection and a dual thread design (range 2 mm), sharper at the apex and gradually lessening toward the neck. Both implants have conical root-shaped geometry and micro-rings at the neck.

All surgeries were performed by postgraduate students of the Implantology Postgraduate Program, FMDUL, Lisbon, Portugal, under the supervision of surgery instructors. Imaging

techniques (panoramic and intraoral radiographs, dental computerized tomographs) were applied for all patients for proper pre-surgical planning.

Immediately before surgery, the mouth was rinsed with a 0.12% chlorhexidine solution (Eludril, Laboratoires Pierre Fabre, France) for 1 minute. The surgical procedures were done under local anesthesia (articaine containing 1:100000 epinephrine – Artinibsa, Inibsa), using loco-regional or infiltrative anesthesia techniques, according to the anatomical region. A midcrestal incision was made, and a full-thickness flap was obtained.

The implants were placed according to the manufacturer's instructions. A one-stage surgical protocol with non-immediate loading was planned for this study. For definition purposes, implants placed in the "anterior" region were meant to replace central and lateral incisors and canines, whereas those in the "posterior" region were placed to replace premolars and molars. Implants were surgically positioned with the implant shoulder at the level of the alveolar bone crest. Primary closure of the wound was made using 3-0 polyglactin 910 single sutures (Vicryl Rapide, Ethicon). Post-operative antibiotic (amoxicillin, 1 g, 12/12 hours for 8 days) and analgesic (paracetamol, 1 g, 8/8 hours for 2 days) were prescribed. All patients were instructed to follow a soft cold diet for the first 24 hours and mouthwash with chlorhexidine 0.2% for a week.

RFA was employed to measure implant stability. RFA measurements were taken immediately after implant placement (T0) and 8 weeks later (T1), using the Osstell Mentor device (Osstell Integration Diagnostics, Goteborg, Sweden). For each implant, ISQ values were measured from the three orientations (buccal, lingual, and occlusal), according to the method described by Rokn.<sup>15</sup> The mean of all measurements was regarded as the final ISQ of the implant. A single operator made all measurements and recordings.

Data were analyzed using IBM SPSS Statistics version 25.0 (IBM Corp., Armonk, NY, USA). The t-test or Mann-Whitney U test and ANOVA or Kruskal-Wallis tests were used according to groups of variables, normality of the distribution, and homogeneity of variance. The t-test was used for groups of two qualitative variables with normal distribution, and the Mann-Whitney U test was applied if they had a non-normal distribution. ANOVA was used for groups of three or more variables if the distribution was normal and variance was homogeneous; if not, Kruskal-Wallis was applied. The level of significance was set at  $p < 0.05$ . Multiple comparisons were made to identify significant differences between groups. The quantitative variables analyzed were mean primary stability and mean secondary stability.

## Results

A total of 45 patients (28 women and 17 men) with an average age of 46 years were included and received 79 implants. No dropouts were registered. The number of implants inserted was 47 (59.5%) in women and 32 (40.5%) in men. The implants' length was 8.0 mm, 10.0 mm, 11.0 mm, and 11.5 mm, and their diameter was 3.75 mm and 4.20 mm. Of the 79 implants studied, 3 MIS Seven implants were excluded due to osseoin-

**Table 1. Implant dimensions and anatomical region for the two groups.**

		C1 Implants (n=40)		Seven Implants (n=39)	
		Cases (n)	Cases (%)	Cases (n)	Cases (%)
<b>Implant diameter</b>	3.75 mm	19	47.5	20	51.3
	4.2 mm	21	52.5	19	48.7
<b>Implant length</b>	8.0 mm	11	27.5	1	2.6
	10.0 mm	14	35.0	19	48.7
	11.0 mm	2	5.0	2	5.1
	11.5 mm	13	32.5	17	43.6
<b>Anatomic region</b>	Anterior maxilla	5	12.5	7	17.9
	Posterior maxilla	5	12.5	10	25.6
	Anterior mandible	1	2.5	1	2.6
	Posterior mandible	29	72.5	21	53.8

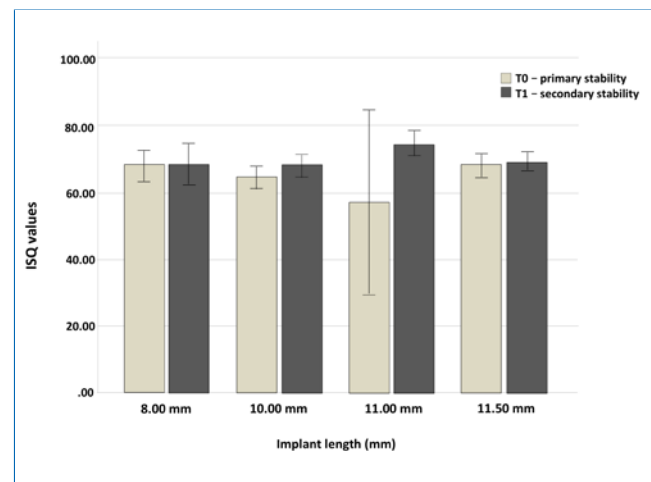
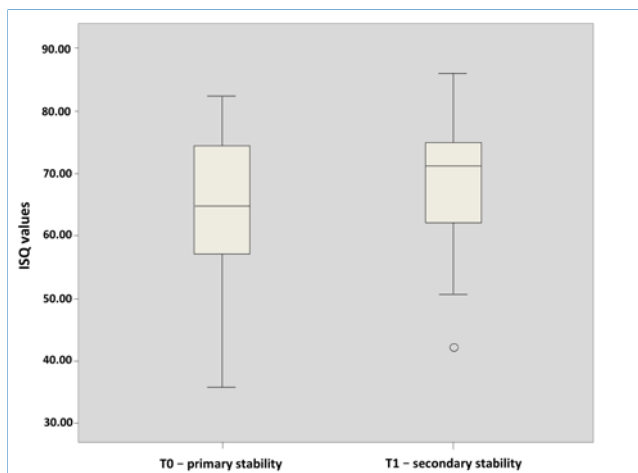
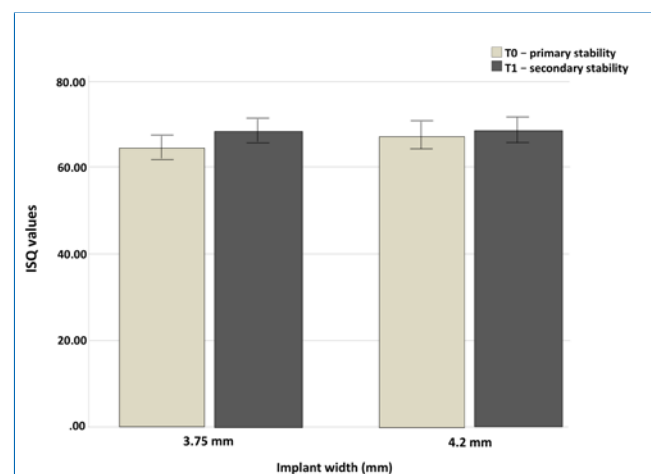
tegration failure, reaching an overall implant survival rate of 96.2%. The anatomical region for implantation, implant types, and implant dimensions are listed in Table 1.

Mean ISQ values were  $65.2 \pm 10.3$  at implant placement and increased statistically significantly after 8 weeks ( $p=0.023$ ), with a mean of  $68.7 \pm 8.6$  (Figure 1). Descriptive statistics of ISQ values distribution regarding implant length and diameter are reported in Figures 2 and 3, respectively.

When considering the overall primary stability (T0) and implant length (8.0 mm, 10.0 mm, 11.0 mm, and 11.5 mm), no statistically significant relationship was observed between length and ISQ values ( $67.9 \pm 7.6$ ,  $63.9 \pm 10$ ,  $57.2 \pm 11.1$ , and  $66.4 \pm 11.3$ ,  $p=0.312$ ). Also, no relationship was found between the implant width (3.75 mm and 4.2 mm) and the ISQ values ( $64.3 \pm 8.7$  and  $66.1 \pm 11.7$ ,  $p=0.445$ ). Regarding implant types, no statistically significant differences were found in the mean primary stability between the two designs of implants studied ( $65.3 \pm 8.8$  for MIS C1 implant and  $65.2 \pm 11.8$  for MIS Seven implant,  $p=0.962$ ).

Analysis of the experimental data also revealed that secondary stability was not affected by implant length or width. No significant differences were observed when comparing the ISQ values for secondary stability between different implant lengths ( $p=0.504$ ). Similarly, there was no correlation between

the implants' diameter and the ISQ values for secondary stability ( $p=0.930$ ). However, different results were obtained when considering the implant design. The secondary stability ISQ

**Figure 2. ISQ values distribution within different implant lengths.****Figure 1. ISQ stability values measured using resonance frequency analysis immediately after implant placement (T0) and 8 weeks later (T1).****Figure 3. ISQ values distribution within different implant diameters.**

**Table 2. RFA mean values and standard deviation (SD) for primary (T0) and secondary stability (T1) regarding implant length, diameter, anatomical region, and implant type.**

		RFA values (T0)		RFA values (T1)	
		Mean	SD	Mean	SD
<b>Implant diameter</b>	3.75 mm	64.3	8.7	68.8	8.2
	4.2 mm	66.1	11.7	68.7	9.1
<b>Implant length</b>	8.0 mm	67.9	7.6	68.4	9.4
	10.0 mm	63.9	10.0	67.9	9.3
	11.0 mm	57.2	11.1	74.7	1.5
	11.5 mm	66.4	11.3	69.2	7.9
<b>Anatomical region</b>	Anterior maxilla	72.8	8.8	75.1	4.4
	Posterior maxilla	63.1	11.3	62.4	8.7
	Posterior mandible	64.6	9.6	69.0	8.3
<b>Implant type</b>	C1	65.3	8.8	67.3	8.1
	Seven	65.2	11.7	70.3	9.0

**Table 3. Scheffe's post-hoc test for multiple comparisons for T0 (primary stability) in relation to the anatomical region.**

(I) Region	(J) Region	Mean Difference (I-J)	Std. Error	Sig.	Confidence Interval (95%)	
					Lower Bound	Upper Bound
<b>Anterior maxilla</b>	Posterior maxilla	9.683*	3.800	0.045	0.190	19.177
	Posterior mandible	8.170*	3.154	0.040	0.290	16.050
<b>Posterior maxilla</b>	Anterior maxilla	-9.683*	3.800	0.045	-19.177	-0.190
	Posterior mandible	-1.513	2.889	0.872	-8.729	5.703
<b>Posterior mandible</b>	Anterior maxilla	-8.170*	3.154	0.040	-16.050	-0.291
	Posterior maxilla	1.513	2.889	0.872	-5.703	8.730

\*The mean difference is significant at the 0.05 level

**Table 4. Kruskal-Wallis post-hoc pairwise comparisons for T1 (secondary stability) in relation to the anatomical region.**

Sample 1 – Sample 2	Test Statistic	Std. Error	Std. Test Statistic	Sig.	Adj. Sig.
<b>Posterior maxilla – Posterior mandible</b>	-14.845	6.704	-2.214	0.027	0.080
<b>Posterior maxilla – Anterior maxilla</b>	35.205	8.603	4.092	0.000	0.000*
<b>Posterior mandible – Anterior mandible</b>	20.361	6.922	2.942	0.003	0.010*

Each row tests the null hypothesis that Sample 1 and Sample 2 distributions are the same. Asymptotic significances (2-sided tests) are displayed.

\*The significance level is 0.05. Significance values have been adjusted by the Bonferroni correction for multiple tests.

values measured for the MIS Seven implant type were significantly higher than those measured for MIS C1 implants ( $p=0.048$ ). ISQ values measured for primary and secondary stability related to implant length, diameter, and implant design are listed in Table 2.

In respect to the anatomical region, implants were distributed in the anterior maxilla (15.2%), posterior maxilla (19.1%), anterior mandible (2.6%), and posterior mandible (63.1%). According to the data obtained, there was a statistical difference between anatomical regions of implantation and primary and secondary stability ( $p<0.05$ ). However, no conclusions could be drawn when making multiple comparisons due to insufficient data from the anterior mandible; thus, these data were not considered. The ISQ values for both primary stability and sec-

ondary stability at the T0 and T1 measurements were significantly higher in the anterior maxilla than in posterior regions of the maxilla and mandible ( $p=0.045$  and  $p=0.040$ , respectively). The significance of differences between anatomical regions is reported in Tables 3 and 4.

## Discussion

This study aimed to determine how the implant design, length, diameter, and anatomical region affect the primary and secondary stability measured by RFA. Our findings suggest that the length and diameter of the implants studied did not influence their stability. Moreover, implant design may



influence secondary stability, whereas intraoral location has a relevant effect on primary and secondary stability.

The effect of implant length on primary stability and implant's long-term prognosis has been a controversial issue. Some studies showed that short implants failed more often than longer implants, while others reported that implant length did not significantly influence implant survival rates.<sup>(16)</sup> Moreover, some studies suggested that increasing implant length plays an important role in decreasing bone stress and increasing implant stability in poor quality bone areas.<sup>17</sup> Likewise, some authors reported that implant diameter influenced ISQ values,<sup>18</sup> whereas others reported no differences.<sup>19,20</sup>

In the present clinical study, the primary and secondary stability were evaluated in two types of implant designs (MIS C1 implants and MIS Seven implants) at implant placement and after a healing period. This study concluded that the implants' length was not significant for primary or secondary stability. These conclusions are in agreement with previous research that found no correlation between implant length and stability.<sup>21</sup> Regarding implant diameter, the results also showed no statistical difference on ISQ values between the different diameters for both primary and secondary stability. These data agree with prior studies that found no correlation between implant diameter and implant stability.<sup>22</sup>

The perception that longer and larger implants have greater stability may not be accurate. Implant design and the surrounding bone may have a more significant effect on implant stability. The implant body design and the thread geometry are significant for improving the mechanical implant stability.<sup>23</sup> Within the limitations of this study, the implant design appears to play a role in implant stability measured after a healing period.

In respect to the bone region, the present study observed that implants placed in the anterior region had more primary stability than implants placed in the posterior regions of the maxilla or mandible. The secondary stability was also poorer in the posterior areas of the jaws. The reason for this may be related to low bone density, which causes decreased implant stability. The poorer implant stability in the posterior area might explain the lower implant success rates reported in the posterior maxilla than in the other regions.<sup>24</sup> In the present study, the three implants that failed were placed in posterior regions of the jaws. The posterior maxilla's reduced bone density and the challenges of mandibular posterior blood supply may explain these failures.

These results agree with those of a previous investigation that found that implants placed in the anterior region had lower failure rates than implants placed in the jaws' posterior region.<sup>25</sup> In a more recent study, no differences were found in ISQs regarding bone types, except between D2 and D3, according to Misch's classification.<sup>26</sup> Some authors stated that, in cases of low bone quality, an optimum increase in implant length and diameter should be taken into account to achieve higher primary stability.<sup>27</sup>

Limited data is available concerning the influence of implant design on implant stability. A randomized clinical trial found a small initial advantage of conical implants with wide pitch compared to semiconical implants and narrow pitch.<sup>28</sup> However, after 90 days, both implant designs showed similar

stability. In the present study, the implant design significantly influenced the ISQ values on secondary rather than primary stability.

Besides the implant geometry, length, or diameter, other parameters have to be considered regarding implant stability and success of dental implants, namely the bone quality, the surgical technique, or the insertion torque. The results of the present study should be confirmed with a larger number of implants, in bone substrates of different densities and with different implant designs.

More long-term controlled clinical trials are needed to confirm the outcomes of the present study, with more patients and fixtures inserted and longer follow-ups, to formulate clear guidelines for implant stability and the success of dental implants.

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## Conclusions

Within the limitations of this study, it can be concluded that, rather than the diameter or length of implants, the anatomical region of implantation has a relevant effect on both primary and secondary stability of dental implants.

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## Ethical disclosures

**Protection of human and animal subjects.** The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

**Confidentiality of data.** The authors declare that they have followed their work center protocols on access to patient data and for its publication.

**Right to privacy and informed consent.** The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

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## Conflict of interest

The authors have no conflicts of interest to declare.

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