



The International Journal of Periodontics & Restorative Dentistry

Official Journal of the Academy of Osseointegration

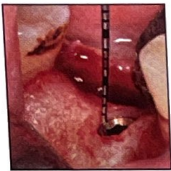


2021/3

Volume 41
Issue 3 • May/June 2021

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Influence of Implant Placement Depth and Soft Tissue Thickness on Crestal Bone Stability Around Implants With and Without Platform Switching: A Comparative Clinical Trial



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This case control study measured early crestal bone changes around subcrestally placed platform-switched implants surrounded by thin soft tissue and compared them with regular, matching-platform implants placed in a supracrestal position and surrounded by thick soft tissue. Sixty-six patients received two-piece internal hex dental implants. Control group patients (n = 33) received implants that had a horizontally matching implant-abutment connection and were placed approximately 0.5 to 1 mm supracrestally. Test group patients (n = 33) received platform-switched implants that were placed about 1.5 mm subcrestally. Clinical examinations were conducted, intraoral radiographs were taken, and statistical analysis was performed. After 2 months, the mean bone loss was 0.2 mm (SD: 0.22 mm; range: 0.1 to 1.2 mm) in the control group and -0.69 mm (SD: 0.65 mm; range: 0 to 2.6 mm) in the test group; this difference was found to be statistically significant ($P < .05$). After 1 year, mean bone loss was 0.28 mm (SD: 0.36 mm; range: 0.1 to 1.63 mm) in the control group and -0.6 mm (SD: 0.55 mm; range: 0.05 to 1.8 mm) in the test group. Platform-switched implants placed in a subcrestal position in vertically thin soft tissues showed statistically significantly more bone loss than non-platform-switched implants placed supracrestally with vertically thick tissues. Int J Periodontics Restorative Dent 2021;41:347–355. doi: 10.11607/prd.5256

Stability of the crestal bone around dental implants is a major concern in dental professions because it is a key factor for treatment success. For decades, dentists focused on alveolar bone qualities and basic surgical principles as main aspects to achieve good outcome.¹ However, guidelines regarding the relation of vertical soft tissue thickness and depth of the implant placement were not covered. Further, it was clinically observed that resorption of the crestal bone is better prevented when tissues surrounding the implants are thicker.^{2,3} Bone is protected from bacteria in the oral cavity by biologic width formation.⁴ The concept of biologic width was first used by Gargiulo et al, who described the dimensions of the dentogingival junction around human teeth.⁵ It has been hypothesized that a similar association might exist between implants as well, and some changes in this relationship may be one of the reasons for early crestal bone loss. This theory first was proved by a canine study wherein thin vertical soft tissues caused crestal bone loss around implants, which established sufficient protection from oral bacteria.⁶ Later clinical studies and systematic reviews established that tissue thickness should be evaluated before treatment, as failure to address it may lead to bone resorption.^{2,3,7}

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Submitted June 20, 2020; accepted August 13, 2020.

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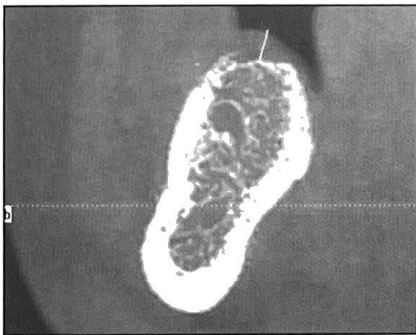


Fig 1 The blue line represents the vertical soft tissue measurement on the CBCT scan.

One way to solve marginal bone loss related to thin tissues is to place the implant subcrestally. This method was known for many years but was first scientifically evaluated by Vervaeke et al in clinical split-mouth comparative study, which showed only 0.04 mm of bone loss in subcrestal implants compared to 0.6 mm in the control epicrestal group.⁸ Although some data exists, submerging implants deep below bone level is still not well-researched, and it is not clear how bone would react to placement of implants with different designs. Another factor influencing marginal bone loss is the microgap. It was shown by in vitro studies that there is a bacterial leakage and colonization of bacteria along the whole connection of implant, starting from the implant-abutment interface.⁹ The microgap is also responsible for formation of inflammatory cell infiltrate in surrounding tissues.^{10,11} It is evident that clinicians should take into consideration two main factors of crestal bone loss: vertical soft tissue thickness and the microgap.

Therefore, the present study aimed to measure early crestal bone

changes around subcrestally placed platform-switched implants with an internal connection of 45 degrees surrounded by thin soft tissue, and to compare the results with those of control matching-platform implants, placed supracrestally and surrounded by thick soft tissue. The null hypothesis was that there would be no difference between the groups.

Materials and Methods

Patients

The subjects who participated in this study were partially edentulous patients of Vilnius University Zalgiris Clinic, Lithuania. The trial's protocol was approved by Lithuanian Bioethics Committee (no Nr.158200-14-752-270). This study took place from 2015 to 2020 and was conducted according to the principles of the Helsinki Declaration. The inclusion criteria were as follows: (1) partially edentulous mandible in the premolar or molar region; (2) age \geq 18 years; (3) no medical contraindication for implant surgery; (4) sufficient alveolar ridge width ($>$ 7 mm) for placement of a 4.6-mm-diameter implant; (5) healed bone sites (at least 6 months after tooth extraction); and (6) no need for bone augmentation procedures. Patients were excluded from study if they fulfilled any of following criteria: (1) poor oral hygiene; (2) suffering from periodontitis; (3) inability to attend follow-up visits; (4) problematic substance users (smoking, alcohol, etc); (5) implant sites were in need of bone augmentation; (6) primary

stability of dental implant $<$ 35 Ncm during surgery.

Soft Tissue Measurement, Group Allocation, and Surgery

Soft tissue thickness of the implant placement site was measured prior to surgery via CBCT scan, which was done with standard double cheek retractors to separate the soft tissue contour (Fig 1). Depending on vertical soft tissue thickness, patients were divided into either the test group (tissue thickness $<$ 2.5 mm) and or the control group (tissue thickness \geq 2.5 mm). Surgical procedures were completed by the same surgeon (S.Z.).

After preparing the operating field and injecting local anesthetic (Ubistesin Forte, 3M ESPE), an incision was made at the center of the edentulous ridge. A buccal full-thickness flap was raised with a periosteal elevator, and the vertical soft tissue thickness of the lingual flap was measured using a periodontal probe (Hu-Friedy). After accurate evaluation of vertical soft tissue thickness, the lingual flap was raised, and the surgeon proceeded with implant placement. Patients in the control group received implants with a horizontally matching implant-abutment connection (4.6-mm diameter; Tapered Internal, BioHorizons), which were placed approximately 0.5 to 1 mm supracrestally. Patients in the test group received platform-switched implants (Tapered Plus, BioHorizons), which were placed about 1.5 mm subcrestally (Fig 2). The accuracy of the

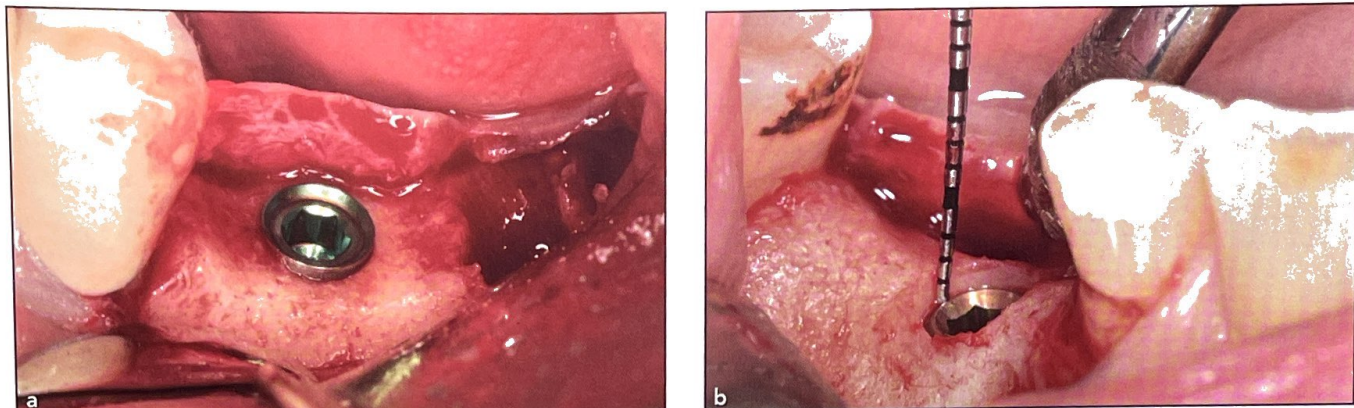


Fig 2 (a) Control group patients had implants placed in a supercrestal position, and (b) test group patients had implants placed in a subcrestal position.

placement was assured by preparing the osteotomy with a drill (Bio-Horizons) longer than the implant (for example, a 12-mm drill length for a 10.5-mm-long implant). After implant placement, straight emergence profile healing abutments were connected, and the soft tissue was sutured without tension using 5/0 interrupted sutures (Vicryl, Ethicon, Johnson & Johnson).

After surgery, patients were recommended to use mouthwash with 0.12% Eludril (Pierre Fabre) twice a day for 1 week. All patients were prescribed postsurgical antibiotics (1 g amoxicillin; Ospamox; Biochemie) twice daily for 7 days and ibuprofen as needed (400 mg; Ibuprom; US Pharmacia). After 1 week postoperative, sutures were removed. During the healing phase, patients were instructed to use a soft toothbrush to clean the healing abutments.

Prosthetic Restoration

Screw-retained fixed restorations were delivered 2 to 3 months af-

ter implant installation. A vinyl-polysiloxane impression material (Express, 3M ESPE) was used for a single-step impression with an individual impression tray. All fixed restorations were radiographically checked for passive fit. After delivery, all patients received dental hygiene instruction using interdental brushes (Curaprox, Curaden).

Clinical Examination

Clinical examination was performed at the 1-year follow-up. During the visit, plaque and bleeding scores were evaluated using the modified Plaque and Bleeding Index, and probing pocket depths were measured.¹²

Radiographic Evaluation

Radiographic evaluation and measurements were completed 2 months after surgery, after prosthesis delivery, and 1 year after prosthesis delivery (Figs 3 and 4). All intraoral radiographs were taken

using a Rinn-like film holder and paralleling technique. Parallelism of radiographs was evaluated before proceeding with measurements, and clear visibility of the implant-abutment interface as a "line" was considered a necessary factor. The radiographs were calibrated by setting 4.6 mm as the diameter of the implant. Calibration and measurements were done with ImageJ software (National Institutes of Health). Bone levels were determined as the distance from the implant-abutment interface, which was used as a reference point. Radiographs were analyzed by one independent examiner.

Statistical Analysis

Statistical analysis was performed using Statistical Analysis System package (version 9.2; SAS). Descriptive statistics were used to describe distributions of variables. Z transformation was performed for quantitative data due to non-normal distribution of data. Student t test was used to evaluate the differences

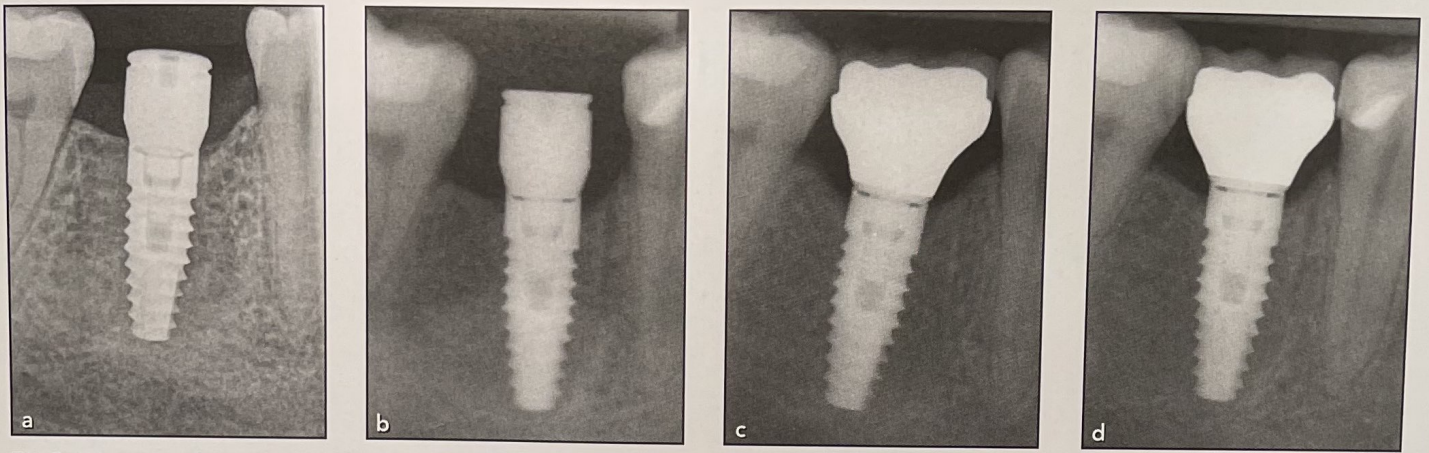


Fig 3 Intraoral radiographs of a control group implant (a) after placement, (b) after 2 months, (c) after prosthesis delivery, and (d) at 1 year.

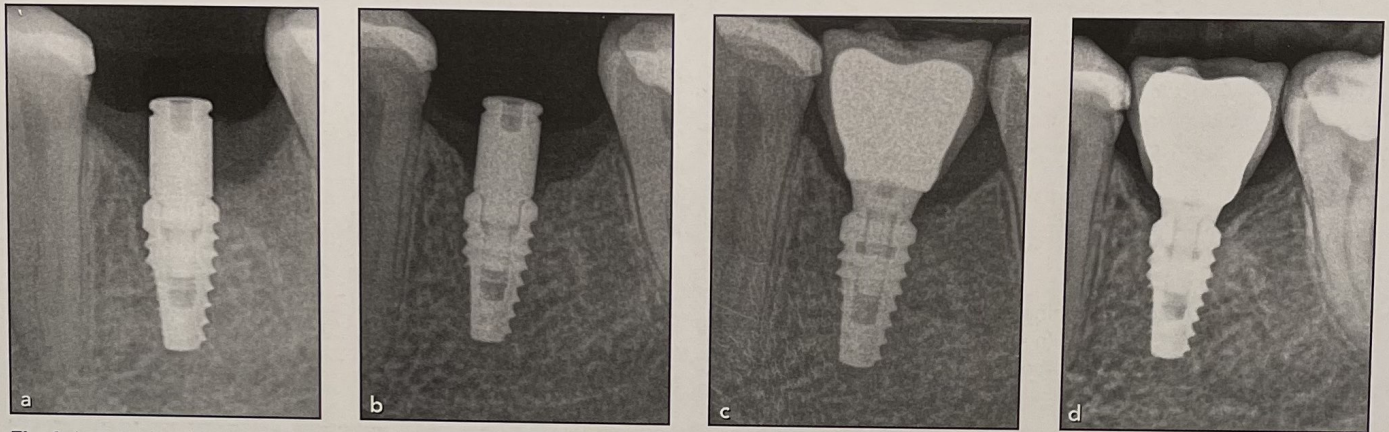


Fig 4 Intraoral radiographs of a test group implant (a) after placement, (b) after 2 months, (c) after prosthesis delivery, and (d) at 1 year.

between the two independent groups with z-transformed data. Two-way analysis of variance with fixed variables was used to evaluate the differences between factors and their interaction. Wilcoxon signed rank sum test was used to evaluate the difference in dependent variables. A two-tailed P value $< .05$ was considered significant with a confidence interval of 95%.

Results

Initially, 70 patients (43 women, 27 men; age range: 23 to 55 years) par-

ticipated in this clinical study. Four patients were excluded during the trial (after implant placement): 1 woman was excluded due to pregnancy and being unable to participate in radiographic evaluation of the results, and 3 patients (2 women, 1 man) were excluded due to lack of cooperation and inability to participate in follow-up visits (Fig 5). The final patient sample included 66 patients (40 women, 26 men) who each received one two-piece internal hex dental implant ($n = 33$ for both control and test groups). The four patients who were excluded from the study were also excluded

from statistical analysis (Table 1). No implants failed during the follow-up, resulting in a 100% survival rate. Statistical analysis revealed a significant difference in bone loss after restoration delivery and at 1 year in the control group ($P < .05$), but no statistically significant difference in bone loss between restoration delivery and the 1-year follow-up in the test group ($P = .397$; Table 2). No statistically significant difference was found between groups and pocket probing depth, Plaque Index, and bleeding on probing (Table 3).

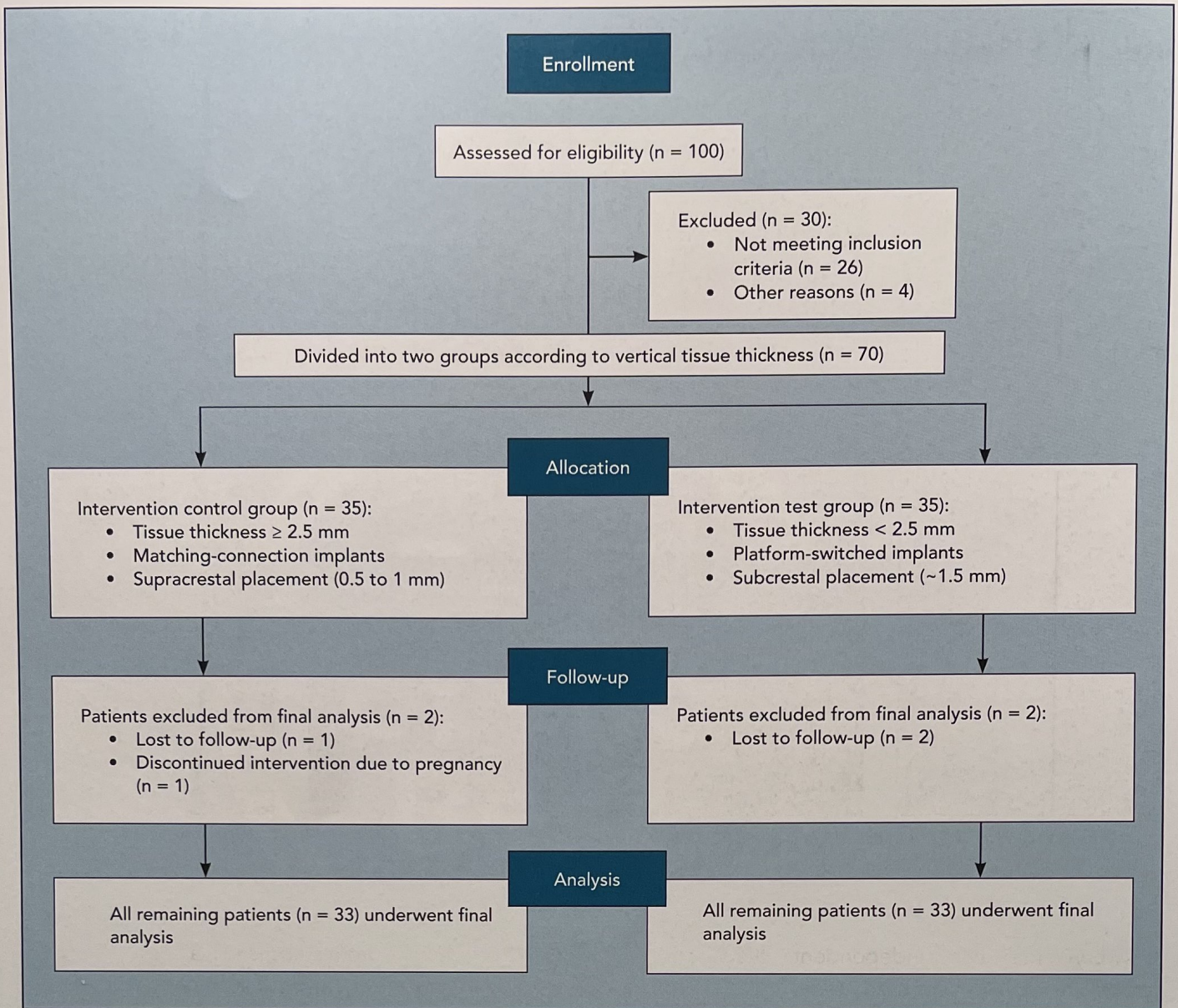


Fig 5 Flowchart of patient enrollment in the study.

Discussion

This study investigated whether implant placement depth in relation with soft tissue thickness has influence on crestal bone stability. The results showed that implants in sites with thick soft tissue showed significantly less bone loss compared to implants in sites with thin soft tissue, even if the implants were in-

stalled subcrestally. Based on this outcome, the null hypothesis that there would be no difference in bone loss between groups was rejected. It is of note that the control group (with soft tissue thickness ≥ 3 mm and non-platform-switched implants with an internal 45-degree connection) showed minor bone loss (0.28 mm) at the 1-year follow-up. A study by van Eekeren et al showed

bone losses of 0.4 mm for bone-level implants and -0.2 mm for tissue-level implants after 1 year of loading.¹³

The results of the control group in the present study can be explained by two major factors: adequate vertical soft tissue thickness and correct implant position, predetermined by the implant design. While tissue thickness was sufficient enough that the protective biologic

Table 1 Crestal Bone Loss and Tissue Thickness in Both Groups After Prosthesis Delivery and After 1 Year of Loading

Parameter	Mean	SD	Minimum	Maximum
Control group (n = 33)				
CBL after prosthesis delivery	0.2	0.22	0.1	1.2
CBL after 1 y	0.28	0.36	0.1	1.63
Tissue thickness	3.04	0.61	2.5	5
Test group (n = 33)				
CBL after prosthesis delivery	0.69	0.65	0	2.76
CBL after 1 y	0.6	0.55	0.05	1.85
Tissue thickness	1.99	0.32	1.4	2.4

CBL = crestal bone loss.
Measurements are shown in millimeters.

Table 2 Comparison of Quantitative Parameters by Wilcoxon Signed Rank Sum Test

Group	Mean (SD) bone loss, mm		P
	Prosthesis delivery	1 y	
Control	0.2 (0.22)	0.28 (0.36)	.003
Test	0.69 (0.65)	0.6 (0.55)	.397

Table 3 Independent Samples t Test Analysis of Periodontal Indices Between Groups

Periodontal index	Control group		Test group		P
	Mean (SD)	Range	Mean (SD)	Range	
Plaque Index	0.09 (0.29)	0–1	0.12 (0.41)	0–1	.65
Bleeding on probing	0.06 (0.24)	0–1	0.09 (0.29)	0–1	.7
Probing pocket depth, mm	2.36 (0.49)	2–3	2.41 (0.67)	2–4	.75

barrier would form with bone loss, implants were placed supracrestally to distance the microgap from the bone level. Already more than a decade ago, Broggin et al described inflammatory cell accumulation in the microgap of the implant-abutment interface around implants with a matching connection, result-

ing in increased crestal bone loss.¹⁴ This means that bone loss can be expected even in sites with a thick tissue biotype if the regular-connection implant is placed crestally, resulting in a microgap position at safe distance from the bone level. Failure to completely understand that crestal bone stability is a multi-

factorial issue may lead to different interpretations of the results. For example, Spinato et al¹⁵ showed no difference in crestal bone stability between thin and thick tissues. However, the implants used in their study had no platform switching and were placed crestally, in which case bone loss is related to too-deep implant

placement rather than tissue thickness.

Many past studies used negative control groups comprised of sites with thin vertical soft tissues to demonstrate that inadequate tissues pose a threat to bone stability.^{2,8,13} However, the present authors believe it is unethical to use those situations as a control any longer, as the literature has shown significant evidence that thin tissues cause bone loss, and ethical committees are reluctant to grant permission for research if the outcome is clearly negative for the patient. Therefore, a positive control group with thick tissue was used in the present study and is suggested for future use.

A separate discussion is needed to understand and explain why subcrestally placed platform-switched implants in thin tissues (test group) had greater bone loss compared to matching-connection implants with a thick tissue type (control group). Generally, it was reported that subcrestal implant placement does not have favorable results with all implant designs; Hermann et al showed that deeper placement of matching-connection implants resulted in more bone loss,¹⁶ while Cochran et al showed favorable results in crestal bone when using platform-switched implants in a subcrestal position.¹⁷ Moreover, Vervaeke et al⁸ also showed almost no bone loss in subcrestally placed platform-switched implants in thin tissues (0.04 mm); this contrasts the outcome of the present study, which showed 0.68 mm of bone loss with platform-switched implants in thin tissues. Both studies used platform

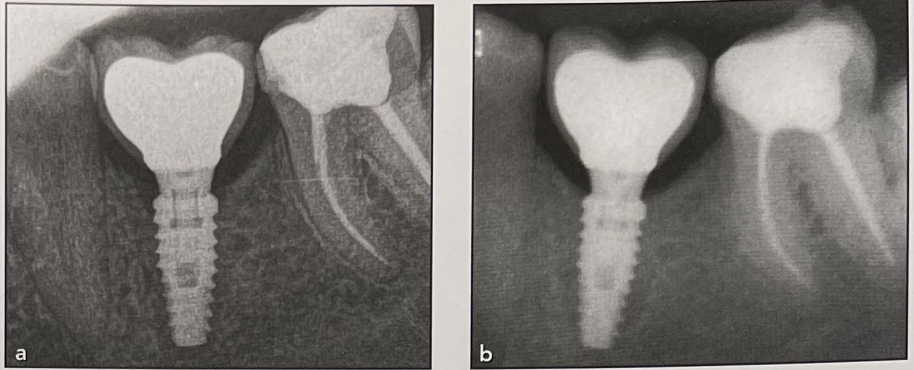


Fig 6 A case of bone remineralization in a test group patient. Intraoral radiographs were taken (a) after prosthesis delivery and (b) at 1 year.

switching as a prerequisite of bone preservation but achieved different outcomes, and the reason might be the differences between connections—or their stability, to be more precise. Vervaeke et al's⁸ implants had a 5-degree conical connection while the present study's implants had a 45-degree conical connection. It was shown that the smaller the angle of the conical connection, the more stable it is.¹⁸ Additionally, it is suggested that the deeper the position of the implant in the bone, the more connection stability is important.¹⁹ The test group used platform-switched implants, but the connection was not stable enough and resulted in more bone loss than expected.

The present study shows that implants without platform switching could maintain stable crestal bone when the surrounding soft tissue is thick. It was interesting to see how platform-switched implants placed subcrestally did not perform better than the control implants

placed supracrestally. This may be explained by the fact that Tapered Internal and Tapered Plus implants have the same internal connection. It was shown that implants placed subcrestally have a stable conical connection, as the internal hex lacks sufficient stability to be positioned subcrestally. The present results show bone loss of 0.69 mm after 2 months in these cases, which were improved (0.6 mm) after 1 year (Fig 6). This phenomenon could be explained by an unstable healing implant-abutment connection during the 2-month healing phase and its consequential bone demineralization. After delivery of the screw-retained prosthesis, the implant-abutment connection was more stable, microbial leakage at the interface was reduced, and remineralized crestal bone was seen in many cases in the test group. This phenomenon of remineralization was explained by Linkevičius et al,¹⁹ who observed bone maturation after eliminating cement remnants in

cement-induced peri-implantitis. Puisys et al presented similar bone behavior in a series of case reports.^{20,21}

It is impossible to conduct the present study without limitations: It could be argued that groups differed from the beginning (thin vs thick tissues) and did not receive equal treatment, as implants with different designs (platform-switched vs matching-platform) were placed in different relations to the bone level (supracrestally vs subcrestally). However, the goal of the study was to compare different protocols of treatment and how to place implants in different tissue thickness. Scientifically, it would be more correct to use the same implant design in both groups, but that would be incorrect clinically. Using implants with a matching connection in the subcrestal (test) group to match the control group would result in extensive bone loss. Therefore, different implant designs were selected for the different treatment protocols being tested.

Conclusions

Within the limitations of this study, it can be concluded that early bone loss around implants is affected by soft tissue thickness and implant design. Platform-switched implants placed in a subcrestal position with vertically thin soft tissues (test group) showed statistically significantly more bone loss than non-platform-switched implants placed supracrestally when tissues were vertically thick (control group). The

test group implants had a moderate crestal bone resorption of 0.69 ± 0.65 mm at 2 months (prosthesis delivery) in spite of subcrestal implant placement. Stabilization of the implant-abutment connections may lead to improvement of the bone stability by an average of 0.1 mm via the process of remineralization.

Acknowledgments

For the following author contributions, an asterisk indicates that the author was the lead person for that task: Conceptualization: S.Z., A.P., and T.L.* Data curation: S.Z.,* P.A., and T.L. Funding acquisition: T.L.* Formal analysis: S.Z.,* E.V., and L.Z. Investigation: S.Z.,* P.A., A.P., and T.L. Methodology: S.Z.,* A.P., E.V., L.Z., and T.L.* Project administration: S.Z.* and T.L.* Resources: S.Z., P.A., and T.L.* Supervision: A.P. and T.L.* Software: S.Z.,* E.V.,* and L.Z. Validation: S.Z., A.P., and T.L.* Visualization: S.Z.* and T.L. Writing (original draft): S.Z.* and T.L. Writing (review and editing): S.Z.,* E.V., L.Z., and T.L.

The authors declare no conflicts of interest.

References

1. Buser D, von Arx T, ten Bruggenkate C, Weingart D. Basic surgical principles with ITI implants. *Clin Oral Implants Res* 2000;11(suppl 1):s59-s68.
2. Linkevičius T, Apse P, Grybauskas S, Puisys A. The influence of soft tissue thickness on crestal bone changes around implants: A 1-year prospective controlled clinical trial. *Int J Oral Maxillofac Implants* 2009;24:712-719.
3. Linkevičius T, Puisys A, Steigmann M, Vindasiute E, Linkeviciene L. Influence of vertical soft tissue thickness on crestal bone changes around implants with platform switching: A comparative clinical study. *Clin Implant Dent Relat Res* 2015;17:1228-1236.
4. Makigusa K. Histologic comparison of biologic width around teeth versus implants: The effect on bone preservation. *International Dentistry* 2009;12:52-58.

5. Gargiulo AW, Wentz FM, Orban B. Dimensions and relations of the dentogingival junction in humans. *J Periodontol* 1961;32:261-267.
6. Berglundh T, Lindhe J. Dimension of the periimplant mucosa biological width revisited. *J Clin Periodontol* 1996;23:971-973.
7. Suárez-López del Amo F, Lin GH, Monje A, Galindo-Moreno P, Wang HL. Influence of soft tissue thickness on peri-implant marginal bone loss: A systematic review and meta-analysis. *J Periodontol* 2016;87:690-699.
8. Vervaeke S, Matthys C, Nassar R, Christiaens V, Cosyn J, De Bruyn H. Adapting the vertical position of implants with a conical connection in relation to soft tissue thickness prevents early implant surface exposure: A 2-year prospective intra-subject comparison. *J Clin Periodontol* 2018;45:605-612.
9. Dibart S, Warbington M, Su MF, Skobe Z. In vitro evaluation of the implant-abutment bacterial seal: The locking taper system. *Int J Oral Maxillofac Implants* 2005;20:732-737.
10. Abrahamsson I, Berglundh T, Sekino S, Lindhe J. Tissue reactions to abutment shift: An experimental study in dogs. *Clin Implant Dent Relat Res* 2003;5:82-88.
11. Ericsson I, Persson LG, Berglundh T, Marinello CP, Lindhe J, Klinge B. Different types of inflammatory reactions in peri-implant soft tissues. *J Clin Periodontol* 1995;22:255-261.
12. Mombelli A, Lang NP. The diagnosis and treatment of peri-implantitis. *Periodontol* 2000 1998;17:63-76.
13. van Eekeren P, Tahmaseb A, Wismeijer D. Crestal bone changes in macrogeometrically similar implants with the implant-abutment connection at the crestal bone level or 2.5 mm above: A prospective randomized clinical trial. *Clin Oral Implants Res* 2016;27:1479-1484.
14. Broggin N, McManus LM, Hermann JS, et al. Peri-implant inflammation defined by the implant-abutment interface. *J Dent Res* 2006;85:473-478.
15. Spinato S, Stacchi C, Lombardi T, Bernardello F, Messina M, Zaffe D. Biological width establishment around dental implants influenced by abutment height, irrespective of vertical mucosal thickness: A cluster randomized controlled trial. *Clin Oral Implants Res* 2019;30:649-659.

16. Hermann JS, Cochran DL, Nummikoski PV, Buser D. Crestal bone changes around titanium implants. A radiographic evaluation of unloaded non-submerged and submerged implants in the canine mandible. *J Periodontol* 1997;68:1117–1130.
17. Cochran DL, Mau LP, Higginbottom FL, et al. Soft and hard tissue histologic dimensions around dental implants in the canine restored with smaller-diameter abutments: A paradigm shift in peri-implant biology. *Int J Oral Maxillofac Implants* 2013;28:494–502.
18. Kofron MD, Carstens M, Fu C, Wen HB. In vitro assessment of connection strength and stability of internal implant-abutment connections. *Clin Biomech (Bristol, Avon)* 2019;65:92–99.
19. Linkevičius T (ed). *Zero Bone Loss Concepts*, ed 1. Chicago: Quintessence, 2019.
20. Puisys A, Auzbikaviciute V, Minkauskaite A, et al. Early crestal bone loss: Is it really loss? *Clin Case Rep* 2019;7:1913–1915.
21. Puisys A, Auzbikaviciute V, Simkunaite-Rizgeliene R, Razukevicius D, Linkevičius R, Linkevičius T. Bone remineralization around dental implants following conservative treatment after peri-implantitis. *Case Rep Dent* 2019;2019:1–6.