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Comparative Analysis of Radiographic Marginal Bone Level Changes in Crestal vs Subcrestal Placement of Implants: A Retrospective Study

Layla Nemat-bakhsh, DDS¹; Sarina Badakhshan, DDS¹; Shunsuke Kamimura, DDS¹; Joseph Kan, DDS, MS¹; Jaime Lozada, DDS¹

¹Advanced Dental Education in Implant Dentistry, Loma Linda University School of Dentistry, Loma Linda, California, USA

Introduction

Implant placement **depth** has been widely recognized as an important factor influencing **marginal bone** preservation and long-term implant success. Despite favorable clinical outcomes with **both** crestal and subcrestal positioning, challenges remain in identifying the optimal vertical placement to **minimize** early bone remodeling. Previous studies¹⁻¹² have reported **conflicting** results regarding the effect of implant depth on marginal bone level changes and peri-implant health.

Null Hypothesis

There is no statistically significant difference in radiographic marginal bone level changes (MBLC) between implants placed **crestally** and those placed **subcrestally** at follow-ups.

Materials and Methods

Records from a retrospective cohort of **Nobel Biocare** and **Straumann** implants placed in healed sites included **480** implants comprising of **201 crestal** and **279 subcrestal** placements. Only cases with radiographs available at implant placement (T0), prosthetic loading (T1), and a minimum one-year follow-up (T2) were included.

A prior **power** analysis was performed to determine the required sample size. An implant was considered to have survived if it remained clinically **functional** at T2.

Standardized radiographic measurements were used to assess marginal bone level change (MBLC) across time points. All measurements were conducted using **ImageJ 1.54**(NIH,Bethesda, MD, USA). The **apical** corner of the implant platform served as the **reference line** (RL). Marginal bone level (MBL) was defined as the distance from **RL** to the most **coronal** bone to implant **contact** on radiographs. Mesial and distal measurements were obtained and **averaged** to calculate MBLC. To compensate for periapical radiographic **distortion**, the correction formula $X = b \cdot c / a$ was applied where **a** = radiographic implant length, **b** = radiographic distance from RL to first bone contact, **c** = actual implant length, and **X** = corrected distance from RL to first bone contact.

Clinical **variables** collected included probing depth, bleeding on probing, plaque index, keratinized mucosa width, and mechanical complications. Data were anonymized and de-identified prior to analysis. Comparative analyses evaluated **MBLC** differences between **crestal** and **subcrestal** placement across **three** intervals: surgery-to-loading, loading-to-latest follow-up, and surgery-to-latest follow-up. Mean differences were reported with **95%** confidence intervals and **p-values**. Correlation analyses assessed relationships among remodeling intervals using Pearson and Spearman coefficients. Statistical significance was set at $\alpha = 0.05$.

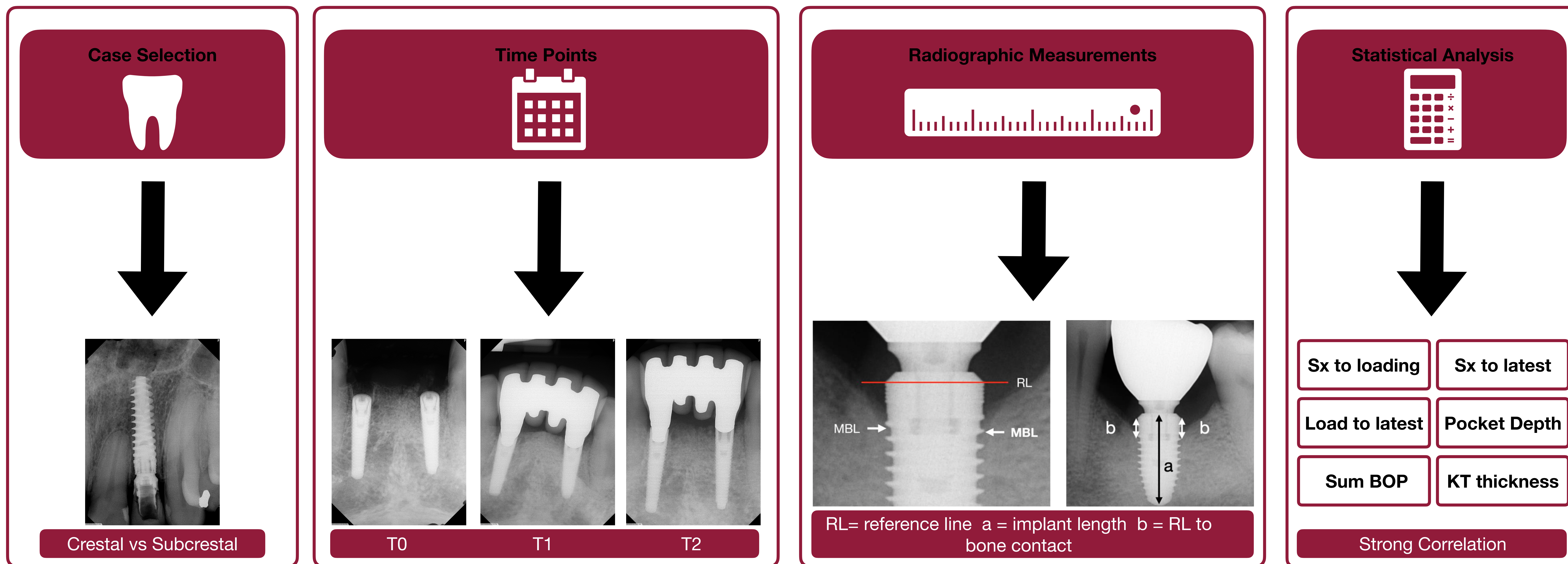


Figure 1. Workflow of retrospective data acquisition and radiographic analysis. Cases were retrospectively selected and categorized by implant placement depth (crestal vs subcrestal). Standardized radiographs were obtained at implant placement (T0), prosthetic loading (T1), and latest follow-up (T2). Marginal bone level changes were measured using a calibrated reference line from the implant platform to the first bone-to-implant contact at mesial and distal aspects. Bone level outcomes were calculated across three intervals: surgery to loading, loading to latest follow-up, and cumulative change. Additional clinical parameters, including probing depth, bleeding on probing, and keratinized tissue thickness, were incorporated into comparative correlation analyses.

Results

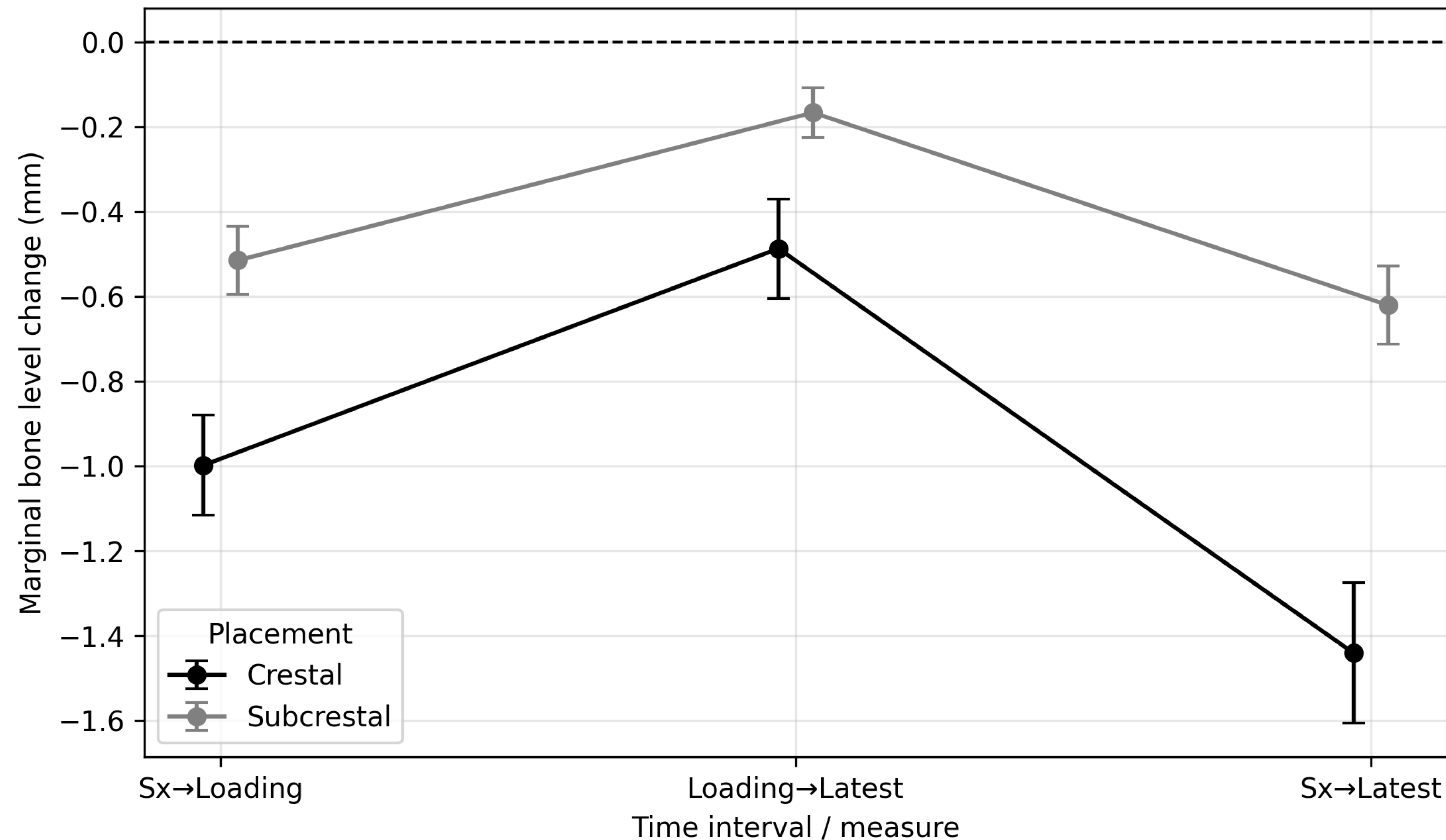


Figure 2. Marginal bone level change (mm) over time demonstrating greater bone preservation in subcrestal compared with crestal implant placement across three intervals: surgery to loading, loading to latest follow-up, and cumulative change. The greatest divergence observed at cumulative follow-up.

Subcrestal implants demonstrated significantly **greater** marginal bone level preservation compared with crestal implants across all time intervals. From time of surgery to loading, the **mean** difference in bone level change was **+0.483 mm** (95% CI: 0.349-0.618; $p < 0.001$). From time of loading to latest follow-up, the difference was **+0.321 mm** (95% CI: 0.164-0.477; $p < 0.001$). Cumulatively (surgery to latest), subcrestal placement showed **+0.821 mm** greater bone preservation (95% CI: 0.162-1.029; $p < 0.01$).

Correlation analyses demonstrated a strong **positive** association between cumulative bone level change and its component intervals ($r = 0.614$ - 0.684 ; $p < 0.001$). A modest **inverse** correlation was observed between **early** and **late** remodeling in subcrestal implants ($r = -0.223$; $p < 0.001$). Increased bleeding on probing was **independently** associated with **greater** bone loss ($\beta = -0.170$ mm per +1 BOP; $p = 0.013$), whereas keratinized tissue thickness, pocket depth, and plaque index were **not significantly** associated with cumulative bone change ($p = 0.21, 0.25, 0.31$). **No significant** interaction was observed between placement depth and bleeding on probing ($p = 0.21$).

Placement	Nobel				Straumann				Location		Failure	Total
Crestal	110				91				Anterior	Posterior	31	201
	Active		Replace		BLT		BLX		45	156		
	39		71		86		5					
	Ti Ultra	Ti Unite	Ti Ultra	Ti Unite	SLA	SL Active	SLA	SL Active				
	19	20	38	33	41	45	0	5				
Placement	Nobel				Straumann				Location		Failure	Total
Subcrestal	166				113				Anterior	Posterior	25	279
	Active		Replace		BLT		BLX		103	176		
	73		93		108		5					
	Ti Ultra	Ti Unite	Ti Ultra	Ti Unite	SLA	SL Active	SLA	SL Active				
	36	37	38	55	63	45	0	5				

Table 1. Distribution of crestal and subcrestal implants by implant system, implant design, surface characteristics, anatomical location, and failure outcomes

Discussion

This **retrospective** analysis evaluated marginal bone level change (MBLC) between **crestal** and **subcrestal** implant placement across early, late, and cumulative healing intervals. Subcrestal implants demonstrated significantly **less** marginal bone loss at all time points, with **greatest** difference observed cumulatively, supporting the influence of placement depth on peri-implant bone stability.¹⁻⁴

Greater divergence over time suggests **sustained** bone preservation beyond **initial** osseointegration. Subcrestal positioning may relocate the implant-abutment interface away from the crestal bone, reducing microgap-associated inflammation and crestal resorption, with

platform switching and stress distribution patterns further contributing to this effect.^{5,6,9,10} Multivariable analysis identified bleeding on probing as **independently** associated with **greater** cumulative bone loss, whereas keratinized tissue thickness was **not** significantly related to bone level change. Correlation findings suggest **early** remodeling patterns may influence **later** crestal stability, consistent with prior reports demonstrating enhanced marginal bone **preservation** with subcrestal placement.^{7,8,11,12}

Conclusion

Within the **limitations** of the retrospective analysis, subcrestal implant placement was associated with significantly **reduced** marginal bone loss compared with crestal positioning across all evaluated intervals. **Increased** bleeding on probing was associated with **greater** bone loss, whereas keratinized tissue thickness was **not** significantly related to bone level changes. These findings suggest that subcrestal placement supports both favorable **initial** remodeling and **sustained** peri-implant bone over time, reinforcing its clinical value for optimizing **long-term** implant stability.



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