

Implant success in patients with a history of chronic periodontitis

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INTRODUCTION

As stated at the 6th European Workshop on Periodontology ¹ more information is needed on the effectiveness of implant therapy based on subjects recruited from private dental clinics. Higher susceptibility for peri-implantitis and peri-implant marginal bone loss was found for periodontal compromised subjects 2,3.



OBJECTIVE

Thus, it was the aim of this retrospective study to analyze implant therapy outcomes in periodontitis susceptible patients of a private periodontal practice.

MATERIAL & METHODS

49 healthy and partially dentate patients (age: 37-78 years) with a history of treated chronic periodontitis and taking part in a maintenance program received a total of 112 bone level placed implants (Brånemark System®). 71 TiUnite® and 41 machined-surface implants were restored with fixed single crowns (Fig. 1-3).







Simultaneous alveolar ridge bone augmentation (BA) using bovine derived xenograft (Bio-Oss® Collagen) in combination with a resorbable collagen membrane (Bio-Guide[®]) was performed in conjunction with 42% of the implants.





Radiographs were taken at insertion, abutment connection and yearly follow-ups (Fig. 1-6). Mean healing period (HP) was 9.5 months (range: 2-26 mths.), mean service period (SP) was 19.5 months (range: 3-59 mths.). Digitized radiographs were assessed by 2 blinded examiners, crestal bone loss was measured on mesial and distal aspects of each implant by ImageJ Software (National Institutes of Health, USA). Implant success was determined as: no pain or tenderness upon function, 0 mobility, no exudate history and radiographic bone loss from initial surgery <2 mm ⁵. Clinical parameters including probing depth (PD), bleeding on probing (BOP), keratinized mucosa width (KM) as well as implant length, implant diameter, crown-to-implant ratio (CIR) and implant surface were analyzed with respect to peri-implant bone loss in a multiple regression model with WALD statistics. Odds ratios (OR) were calculated for peri-implant bone loss (BL) ≥ 2 mm on the basis of implants and of patients.



RESULTS

No implant loss occurred during the observation period (OP). Mean bone loss was 1.51 mm (range: 0.0-4.2 mm / N = 112), 32 implants (28.6 %) showed ≥2.0mm bone loss of which 1.19 mm (range: 0.0-3.7 mm) occurred between insertion and loading. 21 patients (42.8 %) revealed one or more implant(s) with bone loss of \geq 2.0 mm. The overall success rate was 71.4 % (Table 1), with significantly lower success for simultaneously augmented sites (BA) than implants without augmentation procedures (57.6 vs. 81.5 %) and significantly lower success for smokers than for non-smokers (40.9 vs. 80.0 %). Focussed on implant success ⁵, significant odds ratios were calculated for implants: (OR) 5.4 for smoking. (OR) 3.3 for (BA) and (OR) 4.8 for (KM) < 2 mm. On the patients level the odds ratio was (OR) 6.5 in smokers (Table 2).

Table 1: represents the overall success rate of implants with bone loss (BL) <2mm, subdivided in smokers, non-smokers, probing depth (PD) _5mm, <5mm, bleeding on probing (BDP+/) bone augmentation (BA) and no bone augmentation. Prevalence of failure was calculated for patients who exhibited one or more implant(s) with bone loss of 220 mm.

	smoker	non- smoker	PD≥5mm	PD<5mm	BOP +	BOP -	BA	no-BA
Implants:	n=13	n=18	n=9	n=22	n=18	n=7	n=20	n=12
BL>2mm	(59.1 %)	(20.0 %)	(34.6 %)	(25.6 %)	(39.1 %)	(17.1 %)	(42.6 %)	(18.5 %)
Implants:	n=9	n=72	n=17	n=64	n=28	n=34	n=27	n=53
BL<2mm	(40.9 %)	(80.0 %)	(65.4 %)	(74.4 %)	(60.9 %)	(82.9 %)	(57.6 %)	(81.5 %)
Subjects:	n=7	n=14	n=7	n=11	n=13	n=5	n=16	n=5
BL>2mm	(77.8 %)	(35.0 %)	(41.2 %)	(34.4 %)	(48.1 %)	(41.7 %)	(57.1 %)	(23.8 %)
Subjects:	n=2	n=26	n=10	n=21	n=14	n=7	n=12	n=16
BL<2mm	(22.2 %)	(65.0 %)	(58.8 %)	(65.6 %)	(51.9 %)	(58.3 %)	(42.9 %)	(76.2 %)

Table 2: odds ratios calculated for bone loss (BL) \ge 2.0 mm on basis of implants and on patients for local and systemic factors:
OP: observation period, CIB: crown-to-implant ratio, BA: bone augmentation, KM: keratinized mucosa width, IS: implant surface

Odds ratios implant								
parameter	estimate	odds ratio	confidence	e interval	p-value			
age (years)	-0.0176	0.9826	0.9356	1.0318	0.4806			
gender (male vs. female)	-0.0251	0.9752	0.3678	2.5855	0.9598			
OP (days)	0.0003	1.0003	0.9996	1.0010	0.4239			
smoking (smoker vs. non-smoker)	1.6860	5.3966	1.7746	16.4204	0.0030			
CIR (mm)	-0.0228	0.9774	0.1801	5.3039	0.9789			
IS (TiUnite [®] vs. machined)	0.1358	1.1455	0.4921	2.6664	0.7528			
localisation (mandible vs. maxilla)	0.1001	1.1053	0.4229	2.8884	0.8382			
BA (BA vs. no-BA)	1.1853	3.2716	1.2825	8.3457	0.0131			
KM (<2mm vs. ≥2mm)	1.5656	4.7857	1.3304	17.2151	0.0165			

Odds ratios subject								
parameter	estimate	odds ratio	confidence	p-value				
age (years)	-0.0160	0.9842	0.9290	1.0425	0.5869			
gender (male vs. female)	0.5869	1.7000	0.5416	5.3364	0.3633			
OP (days)	0.0001	1.0001	0.9992	1.0010	0.8464			
smoking (smoker vs. non-smoker)	1.8718	6.5020	1.1868	35.5872	0.0310			

REFERENCES

1: Lindhe & Meyle (2008) J Clin Periodontol: (Suppl. 8) 35: 282-285 2: Schou et al. (2006) Clin Oral Imp Res; (Suppl. 2) 17: 104-123 3: Van der Weijden et al. (2005) J Clin. Periodontol: 32(5): 506-511 4: Fransson et al. (2005) Clin Oral Implants Res;16(4):440-6 5: Misch et al. (2008) Implant Dent; 17(1):5-15.

Generalized linear models were used to analyze various risk factors for the outcome of the implants, using a logistic or linear link depending on the type of the outcome variable. Generalized estimation equations were used to estimate the unknown correlation between implants in one patient for this analysis. For patient risk assessment, a patient case was considered as failure if at least one of the implants of the patient failed. The analysis then was done by simple logistic regression. All analyses were performed using SAS 9.2.

Table 3: p-values calculated for local and systemic factors leading to bone loss; Significance level: p < 0.05 HP: healing period, SP: service period, OP: observation period, BA: bone augmentation, KM: keratinized mucosa width, CIR: crown-to-implant ratio, B: implant surace								
	gender	age	time	smoking	BA	КМ	CIR	IS
HP	p=0.74	p=0.15	p=0.001	p=0.03	p=0.0001	p=0.25	-	p=0.48
SP	p=0.39	p=0.64	p=0.004	p=0.75	p=0.93	p=0.61	p=0.46	p=0.23
OP	p=0.96	p=0.29	p=0.049	p=0.036	p=0.0002	p=0.20	p=0.29	p=0.90

Regarding bone loss, a higher incidence existed for implants in smokers and for simultaneously augmented sites (BA) especially during the healing period (HP) (Fig. 7-10). A further significant factor for bone loss was loading time (Table 3).



CONCLUSION

This study indicates that implant success in periodontitis susceptible individuals might be compromised. Implants in smokers revealed more bone loss than in non-smoking patients with periodontitis history. Sites with bone mineral augmentation at implant placement time showed higher amounts of peri-implant bone loss than non-augmented sites during the healing period.

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no-BA

smoke

-0.92 -0.55 0.2500

0.2125 1.90

1.28

1.90

1.9

1.6

19

min media