

Hydroxyapatite-coated implant: Clinical prognosis assessment via a retrospective follow-up study for the average of 3 years

Jun-Hong Jung¹, Sang-Yun Kim², Yang-Jin Yi^{3,4}, Bu-Kyu Lee⁵, Young-Kyun Kim^{2,3*}

¹School of Dentistry, The Graduate School, Seoul National University, Seoul, Republic of Korea

PURPOSE. This research evaluated clinical outcomes of two types of hydroxyapatite (HA)-coated implants: OT (Osstem TS III-HA, Osstem implant Co., Busan, Korea) and ZM (Zimmer TSV-HA, Zimmer dental, Carlsbad, USA). MATERIALS AND METHODS. The research was conducted on 303 implants (89 of OT, 214 of ZM), which were placed from January 16, 2010 to December 20, 2012. The prognosis was evaluated in terms of success rates, survival rates, annual marginal bone loss, and implant stability quotients (ISQ). The samples were classified into immediate, early, conventional, and delayed groups according to the loading time. **RESULTS.** Overall, there were no significant differences between OT and ZM in success rates, survival rates, and annual marginal bone loss, except for the result of secondary stability. OT showed 77.83 ± 8.23 ISQ, which was marginally higher than 76.09 ± 6.90 ISQ of ZM (P<.05). In terms of healing periods, only immediate loading showed statistically significant differences (P<.05). Differences between OT and ZM were observed in terms of two indices, the annual marginal bone loss (0.17 \pm 0.58 mm/year < 0.45 \pm 0.80 mm/year) and secondary stability (84.36 \pm 3.80 $ISQ > 82.48 \pm 3.69$ ISQ) (P<.05). OT and ZM did not have any statistically significant differences in early, conventional, and delayed loading (P>.05). **CONCLUSION.** OT (97.75%) and ZM (98.50%) showed relatively good outcomes in terms of survival rates. In general, OT and ZM did not show statistically significant differences in most indices (P>.05), although OT performed marginally better than ZM in the immediate loading and 1-stage surgery (P<.05). [J Adv Prosthodont 2018;10:85-92]

KEYWORDS: Dental implant; Hydroxyapatite coating; Outcome

Corresponding author:

Young-Kyun Kim

Department of Oral and Maxillofacial Surgery, Section of Dentistry, Seoul National University Bundang Hospital, 82 Gumi-ro 173 beon-gil, Bundang-gu, Seongnam 13620, Republic of Korea Tel. +82317877541: e-mail, kyk0505@snubh.org

Received May 7, 2017 / Last Revision September 20, 2017 / Accepted December 5, 2017

© 2018 The Korean Academy of Prosthodontics

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/3.0) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

This study was supported by the Medical-Dental Convergence Research Program of Seoul National University (860-20140121).

INTRODUCTION

Since the introduction of dental implant treatments in 1981 by Brånemark, efforts to achieve more rapid and stable osseointegration have continued.¹ Researchers have taken efforts to improve osseointegration and osteogenesis by coating the implant surface with hydroxyapatite, a major component of bone.² In particular, with demands for shortening the time of implant treatment and simplification of surgical procedures, especially when dealing with patients with poor bone quality, hydroxyapatite coating received much attention as a solution to address these issues.^{3,4} However, early efforts were plagued by vulnerability to perimplantitis, hydroxyapatite dissolving after surgery, and failed interfacial adhesion between implant and hydroxyapa-

The Journal of Advanced Prosthodontics 85

²Department of Oral and Maxillofacial Surgery, Section of Dentistry, Seoul National University Bundang Hospital, Seongnam, Republic of Korea

³Department of Dentistry & Dental Research Institute, School of Dentistry, Seoul National University, Seoul, Republic of Korea ⁴Department of Prosthodontics, Section of Dentistry, Seoul National University Bundang Hospital, Seongnam, Republic of Korea

⁵Department of Oral and Maxillofacial Surgery, Asan Medical Center, College of Medicine, Ulsan University, Seoul, Republic of Korea

tite. This led to the disappearance of hydroxyapatite from the market.⁵⁻⁷ Subsequently, attention was focused on hydroxyapatite again owing to improved interfacial adhesion between implant and hydroxyapatite and advancements made in processes that allowed a more uniform coating. Examples of such advancements include ion-sputtering, process of thermal composition, and thermal plasma coating.8-11 Each implant manufacturer has its own proprietary coating process. A good example of this is MP-1, which is a thermal plasma coating technology applied to ZM (Zimmer TSV-HA, Zimmer Dental, Carlsbad, CA, USA) implants. The Zimmer TSV-HA used in this study achieved a 97% success rate because it increased the HA crystallization rate to 97% with the application of a plasma coating over the HA and a special MP-1 process using a compressed hydration heat treatment. 12,13 Other manufacturers also produced several thermal plasma coating implants such as OT (Osstem TS III-HA, Osstem Implant Co., Busan, Korea). To improve osseointegration, it is important to increase the crystalline hydroxyapatite content while maintaining the strength of the implant itself. Consequently, manufacturers of hydroxyapatite-coated implants are developing their products with an emphasis on this aspect. OT was treated by the hybrid-type coating with HA and RBM, and the HA crystallinity was higher than 98%. Two millimeters of the upper fixture was RBM surface, and the rest was HA surface to improve the bone response in the cancellous bone, to restrict deposits of bacteria, and to lower marginal bone loss. The Ca/P ratio of the HA surface used in the above experiments was 1.76.

The present study conducted a comparative analysis on the prognosis between two thermal plasma hydroxyapatite-coated implants OT and ZM through a retrospective follow-up study for identifying the mid-term clinical outcomes (maximum 4.52 years, average 2.63 years) of OT, as only few studies have assessed the long-term clinical outcomes. Since the present study examined several implants (n = 303), it also reported on the results that were found for each implant under various conditions such as duration of treatment and surgical technique. The present study conducted a multi-dimensional analysis using success rate, annual marginal bone loss (aMBL), and implant stability quotient (ISQ) as prognostic indices, in addition to survival rates that previous studies primarily examined.

MATERIALS AND METHODS

Between January 2010 and December 2012, 348 hydroxyapatite-coated implants were placed in 185 patients at Section of Dentistry, Seoul National University Bundang Hospital (SNUBH). Among them, 303 implants (89 OT and 214 ZM) that met criteria of the retrospective study were included in the study. The experimental group consisted of 161 patients (87 men and 74 women) with a mean age of 52.65 years (Table 1).

All implant placement procedures were performed by a single oral maxillofacial surgeon, while all prosthetic treat-

ments were performed by a single prosthodontist. All procedures were performed according to the manufacturer's protocol. The present study was conducted under the approval of Institutional Review Board at SNUBH (IRB No: B-1503-290-104). All participants received sufficient explanation on the details of the clinical trial and signed a consent form.

Inclusion criteria of this study were as follows: 1) patients who were 18 years or older 2) patients with manageable systemic disease 3) patients who had undergone minor guided bone regeneration and/or sinus lifting. Exclusion criteria were as follows: 1) patients who withdrew during the middle of the study 2) patients with insufficient medical records and/or radiologic examination records 3) patients who received prosthetic treatment at another hospital after fixture placement.

The implantation sites were as follows: 207 on the maxilla and 96 on the mandible and 34 on the anterior region, 72 on the premolar region, and 197 on the molar region. The highest number of implants was placed on maxillary molar region (129) (Table 2). Placement procedures consisted of two hundred (200) 1-stage surgeries and one hundred and three (103) 2-stage surgeries. The second surgery or impression taking was performed 2 - 3 months and 4 - 6 months after 1st implant placement surgery in mandible and in maxilla, respectively. With respect to classification according to loading types, conventional loading was the most common with 143, followed in order by immediate loading

Table 1. Number of patients and implants

	OT	ZM	Total
Initial number of patients	52	133	185
Mid-term drop out	7	17	24
Final number of patients	45	116	161
Males	26	61	87
Females	19	55	74
Mean age	53.82	51.48	52.65
Total number of implants	89	214	303

Table 2. Number of implants by placement site

	Site	OT	ZM	Total
Maxilla	Anterior	7	14	21
	Premolar	21	36	57
	Molar	39	90	129
Mandible	Anterior	0	13	13
	Premolar	4	11	15
	Molar	18	50	68
Total		89	214	303

with 68, delayed loading with 67, and early loading with 25 (Table 3). Regarding the diameter of implants placed, 5-mm implant was the most common with 60 among OT, while 4.7-mm implant was the most common with 123 among ZM (Table 4). In both groups, 10-mm implant was the most common (OT with 64, ZM with 144) length (Table 5).

The medical records and radiographs of the participants were retrospectively analyzed. Marginal bone loss (MBL), ISQ, success rate, and survival rate were investigated as outcome indices. In addition, comparisons were made accord-

Table 3. Number of implants by surgical and loading methods

	Surgery method			Loadi	ng method	
	1-stage	2-stage	Immediate	Early	Conventional	Delayed
OT	63	26	34	7	31	17
ZM	137	77	34	18	112	50
Total	200	103	68	25	143	67

Table 4. Number of implants by diameter

(#/mm)	3.5	4	4.5	5	3.3	3.7	4.1	4.7	6	Total
ОТ	1	9	19	60						89
ZM					15	15	40	123	21	214

Table 5. Number of implants by length

(#/mm)	8	8.5	10	11.5	13	Total
ОТ		6	64	16	3	89
ZM	42		144	23	5	214

ing to the duration of healing and surgical method. The duration of healing refers to the time until prosthetic loading is exerted on the fixture, which was classified as immediate, early, conventional, and delayed loading. According to 2002 Barcelona consensus, immediate, early, conventional, and delayed loading was established as within 48 hours, 48 hours to ≤ 3 months, ≥ 3 months to ≤ 6 months, and ≥ 6 months, respectively.¹⁴ Surgical method was classified as either 1-stage or 2-stage depending on whether a secondary surgery was required.

Periapical radiographs were acquired during regular follow-up via paralleling technique using RVG 6200 system (Carestream dental, Stuttgart, Germany). Picture archiving and communication system (PACS) used was Infinitt Marosis (Marotech, Seoul, Korea), and this was used to adjust the brightness and contrast of the radiographs to a set level. Marginal bone resorption was analyzed by 2 radiographs, one taken after implant placement and the other from the final examination dividing the changes in the height of marginal bone around the implant by imaging period. On Infinitt software, the height at the highest point of the marginal bone was measured relative to the top of the fixtureabutment connection. For error correction, the value from the software and actual fixture length were multiplied after establishing a proportional expression. Measurements were taken from two points, mesial and distal, and the means of values measured to two decimal places were used. Marginal bone height was analyzed by substituting the fixture height recorded at the time of implant placement into the equation shown in figure 1. Using the method shown below, MBL (in mm) was derived by subtracting the final marginal bone level from the marginal bone level after the prosthesis was attached (Fig. 1 and Fig. 2). Since each case had different time frame on MBL measurement, it was necessary to compensate for such differences. The period from the first measurement date to the final measurement date was recorded in unit of days and 365 days was used as one year to divide

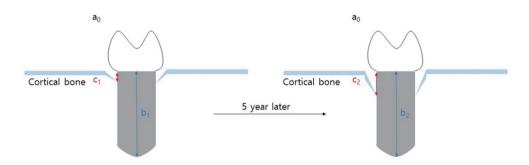


Fig. 1. Marginal bone loss.

Estimated Level of Marginal Bone = Measured Level of Marginal Bone ×

Actual Length of Fixture Measured Length of Fixture

Marginal Bone Loss = 1st Estimated Value - Final Estimated Value

Actual length of fixture: a, Measured length of fixture: b, Measured level of marginal bone: c,

Estimated level of marginal bone: $X = c \times \frac{a}{b}$

Marginal bone loss = $X_2 - X_1$

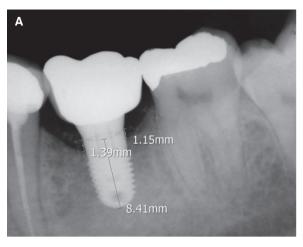




Fig. 2. Example of measurement using periapical radiographs. (A) Left radiograph was taken after delivering a prosthesis, (B) Right radiograph was taken during follow-up period. Both radiographs are ZM implants in the same patient.

the calculated values. This was expressed as aMBL and was compared accordingly. All values were obtained using Excel (Microsoft, Redmond, WA, USA) by setting up the aforementioned function.

Annual marginal bone loss (mm/year)

Implant stability quotient (ISQ) was measured using the Osstell Mentor (Osstell, Gothenburg, Sweden) device. The initial ISQ was measured immediately after implantation, while the final ISQ was measured during secondary surgery for exposing the implant or during acquisition of prosthetic impression when 1-stage surgery was used for implantation. The mean interval between the two measurements was 135.07 days (OT 145.53 days and ZM 132.65 days). To compensate for the differences in measurement periods, Daily ISQ gain (ISQ/day) index was then obtained and compared between groups.

daily ISQ gain (ISQ/day)
$$= \frac{ISQ}{Interval between Initial and Final measure (Days)}$$

Successful implantation was determined according to the definition provided by Albrektsson in 1986, which is the criterion that is most often cited: 1 no persistent pain, discomfort, or dysesthesia 2 no abscess near the implant 3 no tooth mobility 4 no radio-opacity near the implant 5 show MBL of \leq 1 mm for 1 year after prosthetic loading 6 show aMBL of \leq 0.2 mm from 1 year after prosthetic loading 7

show 5-year success rate of \geq 85% and 10-year success rate of \geq 80% 8) the percentage of numbers that satisfy these criteria shall be reported as the success rate.

Cumulative survival rate represented the percentage of implants that were not removed from implant placement to the time of final screening. 16,17

Data were rearranged using Excel (Microsoft, Redmond, WA, USA) and statistical significance was tested using SPSS version 12.0 (SPSS Inc., Chicago, IL, USA). Mann-Whitney test was performed on the outcome values after normality test.

RESULTS

OT and ZM groups did not show statistically significant differences in success rate (P > .05) and survival rate (P > .05). With respect to success rate, OT and ZM groups showed 92.1% and 94.4%, respectively. Both groups showed excellent success rates that were higher than the 5-year success rate criterion given by Albrektsson (85%). 15 Meanwhile, aMBLs in OT and ZM groups were 0.27 \pm 0.89 mm/year and 0.32 ± 0.70 mm/year, respectively, showing a slightly higher value for OT group, but with no statistically significant difference (P > .05). Initial ISQ also showed no statistically significant difference between the two groups (P >.05). The items in which the two implants showed statistically significant difference were final ISQ (P = .03) and daily ISQ gain (P = .05). In the final ISQ item, OT and ZM groups showed values of 77.83 ± 8.23 and 76.09 ± 6.90 , respectively. OT group showed slightly higher value, and as a result, excellent results were seen in daily ISQ gain (0.10 \pm $0.13 > 0.07 \pm 0.14$), which is an indicator of increase in ISQ rate (Table 6).

Table 6. Annual bone loss, implant ISQ, success rate, and survival rate (mean \pm SD)

	OT	ZM
aMBL (mm/year)	0.27 ± 0.89	0.32 ± 0.70
Initial ISQ (ISQ)	69.20 ± 15.00	68.45 ± 11.26
Final ISQ (ISQ)	77.83 ± 8.23*	$76.09 \pm 6.90^*$
Daily ISQ gain (ISQ/day)	$0.10 \pm 0.13^*$	$0.07 \pm 0.14^*$
Success rate (%)	92.13	94.44
Survival rate (%)	97.75	98.15

aMBL: average marginal bone loss, SD: standard deviation, ISQ: implant stability quotient

Results based on classification of 4 types of loading indicated that only the group that had immediate loading applied showed statistically significant different results in aMBL (P = .01) and final ISQ (P = .01). During immediate loading, aMBL was higher in the OT group (0.17 \pm 0.58 mm/year) than in the ZM group (0.45 \pm 0.80 mm/year). Similar to the results of the final ISQ index for the same loading method, OT group was marginally higher with 84.36 \pm 3.80 ISQ, as compared to ZM group with 82.48 \pm 3.69 ISQ. In early, conventional, and delayed loading, OT and ZM groups showed no significant differences (P > .05; Table 7).

Comparison of results based on dividing the surgical method as 1-stage and 2-stage indicated that OT group showed significantly superior aMBL (P = .01) and final ISQ (P = .00) than ZM group for 1-stage surgery. With respect to aMBL index, OT group showed a value of 0.23 ± 0.55 mm/year, which represented less bone resorption by about 0.07 mm/year, as compared to 0.3 \pm 0.54 mm/year in the ZM group. With respect to final ISQ index, OT group showed more stable results with 80.95 ± 6.18 ISQ, as compared to 77.73 ± 6.30 ISQ in the ZM group. When 2-stage surgery was applied, OT and ZM groups did not show significant differences (P > .05; Table 8).

Initial tooth mobility was the most common early complication. Most cases that showed tooth mobility in the OT group resulted in removal of the implant due to failed osseointegration. Contrarily, osseointegration was successful in the ZM group, except for 2 cases. Other than tooth mobility, biological complications included infection, failed osseointegration, paralysis, and fracture (Table 9).

Among late stage complications, prosthesis-related complications were most common in both OT and ZM groups, including prosthesis fall out, prosthesis fracture, screw loosening, and implant structure (fixture or abutment) fracture. Prosthesis-related complications were followed in order by peri-implantitis, bone loss, and gingival recession (Table 10).

Table 7. Results specifically classified by loading methods $(mean \pm SD)$

Loading	OT	ZM
Immediate		
aMBL (mm/year)	$0.17 \pm 0.58^*$	$0.45 \pm 0.80^*$
Initial ISQ (ISQ)	74.96 ± 8.20	77.74 ± 6.22
Final ISQ (ISQ)	84.36 ± 3.80*	$82.48 \pm 3.69^*$
Success rate (%)	82.86	79.41
Survival rate (%)	94.29	100
Early		
aMBL (mm/year)	0.33 ± 0.63	0.21 ± 0.29
Initial ISQ (ISQ)	76.26 ± 11.01	71.39 ± 14.24
Final ISQ (ISQ)	80.31 ± 7.10	75.92 ± 6.00
Success rate (%)	100	94.44
Survival rate (%)	100	100
Conventional		
aMBL (mm/year)	0.22 ± 0.52	0.22 ± 0.46
Initial ISQ (ISQ)	64.07 ± 19.96	67.62 ± 9.44
Final ISQ (ISQ)	74.09 ± 7.37	75.46 ± 6.26
Success rate (%)	96.67	99.12
Survival rate (%)	100	100
Delayed		
aMBL (mm/year)	0.25 ± 0.26	0.5 ± 1.06
Initial ISQ (ISQ)	61.5 ± 11.80	63.3 ± 12.21
Final ISQ (ISQ)	73.22 ± 8.86	73.21 ± 7.60
Success rate (%)	94.12	95.92
Survival rate (%)	100	100

aMBL: average marginal bone loss, SD: standard deviation, ISQ: implant stability quotient

Table 8. Results specifically classified by surgery method $(mean \pm SD)$

	OT	ZM
1-stage		
aMBL (mm/year)	$0.23 \pm 0.55^*$	$0.3 \pm 0.54^*$
Initial ISQ (ISQ)	73.46 ± 9.77	70.78 ± 10.80
Final ISQ (ISQ)	80.95 ± 6.18*	$77.73 \pm 6.30^*$
Success rate (%)	92.06	92.09
Survival rate (%)	96.83	98.56
2-stage		
aMBL (mm/year)	0.19 ± 0.37	0.35 ± 0.91
Initial ISQ (ISQ)	56.43 ± 20.23	64.01 ± 10.84
Final ISQ (ISQ)	71.35 ± 8.31	73.63 ± 7.05
Success rate (%)	92.31	98.70
Survival rate (%)	100	100

^{*} P < .05

^{*} P < .05

^{*} P < .05

Table 9. Early biological complications

Early complication type (patients)	ОТ	ZM
Initial tooth mobility	3	7
Infection	2	3
Failed osseointegration (removal)	2	2
Paralysis	1	1
Fracture	1	0
Total	9	13

Table 10. Delayed complications

Delayed complications (patients)	ОТ	ZM
Prosthetic complications	3	3
Peri-implantitis	2	3
Bone loss	3	2
Gingival recession	0	1
Total	8	9

DISCUSSION

Achieving rapid osseointegration with long-term stability is a prerequisite for satisfying patients who demand immediate restoration after an implant surgery. For the past 20 years, many researchers have attempted to use hydroxyapatite, a bone component, as coating material to shorten the osseointegration process.² Meanwhile, there have been efforts to minimize complications such as peri-implantitis.5-7 As a result, recent studies have reported survival rates of 93.2 -98.5% as clinical outcomes of hydroxyapatite-coated implants.11 Guttenberg reported a 9-year survival rate of 93.3% for Integral (Calcitek Inc., Carlsbad, CA, USA), while Saadoun reported an 8-year survival rate of 96.1% for Steri-Oss (Yorba Linda, CA, USA) implant. Jones reported a 5-year survival rate of 98.5% for Sterngold/implamed (Atteboro, MA, USA) implant. 18-20 In the present study, the survival rates in the OT and ZM groups were 97.75% and 98.15%, respectively. In terms of survival rate, these results indicated that the survival rate in the present study represented superior outcome value than studies that used other hydroxyapatite-coated implants. In terms of success rate, OT and ZM groups showed 92.1% and 94.4%, respectively, which represented excellent success rates that were higher than Albrektsson's 5-year success rate of 85%. 15 Meanwhile, the present study clearly distinguished the results on success rates according to Albrektsson's definition of success rate. It holds significance.

A study by Kim in 2013 that used the same implants as the present study examined the outcomes for 12 months, and demonstrated that the clinical outcome of survival rates for both OT and ZM products was 100%, which exceeded 97.75% and 98.15%, respectively, found in the present study. With respect to MBL, the present study showed greater bone resorption than that observed in previous studies, except for the immediate loading OT group.²¹ It is believed that this outcome may be attributed to the present study having a longer follow-up period. The mean values were derived with inclusion of results from implants that were not removed despite of severe bone resorption. A 2012 paper by Kim reported success rate of 98.1% and 97.9% for OT and ZM, respectively. This study analyzed 1-year follow-up after implants from two manufacturers were placed by applying immediate loading,²² which exceeded success rate of 82.86% and 79.41%, respectively, found in the present study. It is believed that this difference was due to difference in the success criteria applied. The 2012 study by Kim reported a case to be successful if MBL showed < 1.50 mm of bone resorption in the first year, whereas the present study determined success as < 1 mm in the first year and < 0.20 mm per year thereafter.

Comparison of results based on loading methods showed that stable prognosis with 100% survival was found in all cases, except in immediate loading cases. This demonstrated that applying loading within 48 hours after implantation can have a critical impact on prognosis. In the OT group, success rate for early loading was 100%, but the rate decreased to 82.86% with immediate loading. In the ZM group, success rate for early loading was 94.44%, which also dropped rapidly to 79.41% with immediate loading. Meanwhile, such rapid change in prognosis was not found between early and conventional loading or between conventional and delayed loading. Consequently, it is still difficult to actively use immediate loading that has uncertain prognosis in actual clinical settings, and thus, it is desirable to apply early loading as an alternative to satisfy the patient's demand for speedy completion of treatment.

In comparison to other studies that examined immediate loading, a study by Polizzi that used the Brånemark implant (Nobel biocare AB, Sweden) reported immediate implant survival rates of 92.4% and 94.7% in the maxilla and mandible, respectively, while OT and ZM coated with HA used in the present study showed a better prognosis with 94.29% and 100%, respectively.²³

Manufacturers of the existing hydroxyapatite use machined surface of about 1 - 2 mm in the upper area of the fixture to prevent plaque buildup or bacterial infection, but OT has a structural difference of being treated with resorbable blasting media (RBM).²⁴ However, OT and ZM groups in the present study did not show significant differences, and as such, additional studies are needed to determine whether RBM can be used to achieve the same outcome as machined surface.

Comparison between 2-stage and 1-stage surgeries showed that although the surgery itself may be simple, the

additional surgery placed burden on the patient and increased the number of hospital visits. The present study compared the prognosis of implant surgeries performed by an experienced surgeon. Survival rates for OT 2-stage and 1-stage were 100% and 96.83%, respectively. Although 2-stage showed a slightly higher survival rate, prognoses were similar for both methods. Survival rates for ZM 2-stage and 1-stage were 100% and 98.70%, respectively, also showing similar prognoses. This demonstrated that when a patient is burdened by having to undergo another surgery, 1-stage surgery may be a stable and better option.

The data are collected from implants which were placed from 2010 to 2012. Some cases were early failure in osseointegration. Some patients stopped during follow-up. The limitation of this study would be the failure to exclude these data. The loading protocol, prosthetic treatment, occlusal restoration, and follow-up period were variable. Additional studies are needed to further investigate long term prognosis of hydroxyapatite-coating implants with refined data.

CONCLUSION

In a follow-up study up to 5 years, OT and ZM, hydroxyapatite coated implants, did not show statistically significant differences in success rate, survival rate, and aMBL (P > .05). Success rates for OT and ZM were 92.13% and 94.44%, respectively. OT and ZM showed a high survival rate of 97.8% and 98.2%, respectively, demonstrating that hydroxyapatite-coated implants have very high stability in terms of long-term survival rate. In detail, from the perspective of duration of healing or surgical method, OT showed superior aMBL results than ZM for immediate loading and 1-stage surgery (P < .05). Under other conditions, these two implants did not show any significant difference (P > .05). The significance of the present study was that unlike previous studies that reported clinical outcomes by survival rate only, the present study made comparisons with various indices such as success rate, survival rate, MBL, and ISQ.

ORCID

Jun-Hong Jung https://orcid.org/0000-0003-1280-0403 Sang-Yun Kim https://orcid.org/0000-0002-2952-5404 Yang-Jin Yi https://orcid.org/0000-0001-8341-4759 Bu-Kyu Lee https://orcid.org/0000-0001-8888-1719 Young-Kyun Kim https://orcid.org/0000-0002-7268-3870

REFERENCES

- 1. Adell R, Lekholm U, Rockler B, Brånemark PI. A 15-year study of osseointegrated implants in the treatment of the edentulous jaw. Int J Oral Surg 1981;10:387-416.
- 2. de Groot K, Geesink R, Klein CP, Serekian P. Plasma sprayed coatings of hydroxylapatite. J Biomed Mater Res 1987;21: 1375-81.
- 3. Szmukler-Moncler S, Piattelli A, Favero GA, Dubruille JH. Considerations preliminary to the application of early and im-

- mediate loading protocols in dental implantology. Clin Oral Implants Res 2000;11:12-25.
- 4. Gapski R, Wang HL, Mascarenhas P, Lang NP. Critical review of immediate implant loading. Clin Oral Implants Res 2003; 14:515-27.
- 5. Whitehead RY, Lucas LC, Lacefield WR. The effect of dissolution on plasma sprayed hydroxylapatite coatings on titanium. Clin Mater 1993;12:31-9.
- 6. Wheeler SL. Eight-year clinical retrospective study of titanium plasma-sprayed and hydroxyapatite-coated cylinder implants. Int J Oral Maxillofac Implants 1996;11:340-50.
- 7. Zablotsky MH. Hydroxyapatite coatings in implant dentistry. Implant Dent 1992;1:253-7.
- 8. Overgaard S, Søballe K, Josephsen K, Hansen ES, Bünger C. Role of different loading conditions on resorption of hydroxyapatite coating evaluated by histomorphometric and stereological methods. J Orthop Res 1996;14:888-94.
- 9. Yoshinari M, Ozeki K, Sumii T. Properties of hydroxyapatitecoated Ti-6Al-4V alloy produced by the ion-plating method. Bull Tokyo Dent Coll 1991;32:147-56.
- 10. Sugiyama T, Miake Y, Yajima Y, Yamamoto K, Sakurai K. Surface observation of thin hydroxyapatite-coated implants at 80 months after insertion. J Oral Implantol 2011;37:273-8.
- 11. Lee JJ, Rouhfar L, Beirne OR. Survival of hydroxyapatitecoated implants: a meta-analytic review. J Oral Maxillofac Surg 2000;58:1372-9.
- 12. Kim SS, Lee JH, Yu SH, Lee HJ, Moon JW, Park IS, Sohn DS. Comparative study of removal torque of 3 different hydroxyapatite coated implants in the femur of rabbits. J Korean Assoc Oral Maxillofac Surg 2011;37:49-53.
- 13. Eom TG, Kim JH, Cho IH, Jeong CM, Cho YS, Kim YK. Experimental study about the bony healing of hydroxyapatite coating implants. J Korean Assoc Oral Maxillofac Surg 2011;
- 14. Aparicio C, Rangert B, Sennerby L. Immediate/early loading of dental implants: a report from the Sociedad Española de Implantes World Congress consensus meeting in Barcelona, Spain, 2002. Clin Implant Dent Relat Res 2003;5:57-60.
- 15. Albrektsson T, Zarb G, Worthington P, Eriksson AR. The long-term efficacy of currently used dental implants: a review and proposed criteria of success. Int J Oral Maxillofac Implants 1986;1:11-25.
- 16. Naert I, Koutsikakis G, Duyck J, Quirynen M, Jacobs R, van Steenberghe D. Biologic outcome of implant-supported restorations in the treatment of partial edentulism. part I: a longitudinal clinical evaluation. Clin Oral Implants Res 2002;13: 381-9.
- 17. O'Roark WL. Survival rate of dental implants: an individual practitioner's anecdotal review of 25 years of experience. J Oral Implantol 1997;23:90-103.
- 18. Guttenberg SA. Longitudinal report on hydroxyapatite-coated implants and advanced surgical techniques in a private practice. Compend 1993:S549-53.
- 19. Saadoun AP, Le Gall MG. An 8-year compilation of clinical results obtained with Steri-Oss endosseous implants. Compend Contin Educ Dent 1996;17:669-74, 676.
- 20. Jones JD, Lupori J, Van Sickels JE, Gardner W. A 5-year com-

- parison of hydroxyapatite-coated titanium plasma-sprayed and titanium plasma-sprayed cylinder dental implants. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 1999;87:649-52
- 21. Kim YK, Ahn KJ, Yun PY, Kim M, Yang HS, Yi YJ, Bae JH. Effect of loading time on marginal bone loss around hydroxyapatite-coated implants. J Korean Assoc Oral Maxillofac Surg 2013;39:161-7.
- 22. Kim YK, Lee JH, Lee JY, Yi YJ. A randomized controlled clinical trial of two types of tapered implants on immediate loading in the posterior maxilla and mandible. Int J Oral Maxillofac Implants 2012;28:1602-11.
- 23. Polizzi G, Grunder U, Goené R, Hatano N, Henry P, Jackson WJ, Kawamura K, Renouard F, Rosenberg R, Triplett G, Werbitt M, Lithner B. Immediate and delayed implant placement into extraction sockets: a 5-year report. Clin Implant Dent Relat Res 2000;2:93-9.
- 24. Morris HF, Ochi S. Hydroxyapatite-coated implants: a case for their use. J Oral Maxillofac Surg 1998;56:1303-11.