

The seven-year cumulative survival rate of Osstem implants

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Abstract (J Korean Assoc Oral Maxillofac Surg 2014;40:68-75)

Objectives: This study was performed to analyze the cumulative survival rate of Osstem implants (Osstem Implant Co., Ltd.) over a seven-year period.

Materials and Methods: A total of 105 patients who had 467 Osstem implants that were placed at the Section of Dentistry, Seoul National University Bundang Hospital (Seongnam, Korea) from June 2003 through December 2005 were analyzed. The life table method and a cross-tubulation analysis, log rank test were used to evaluate the survival curve and the influence that the prognostic factors. The prognostic factors, i.e., age and gender of patients, diameter and length, type of implants, bone graft history and loading time were determined with a Cox proportional hazard model based on logistic regression analysis.

Results: The seven-year cumulative survival rate of Osstem implants was 95.37%. The Cox proportional hazard model revealed that the following factors had a significant influence on survival rate; increased diameter, reduced prosthetic loading period and performance of bone grafting.

Conclusion: The osstem implants showed satisfactory results over the seven-year study period.

Key words: Dental implants, Survival rate

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I. Introduction

Albrektsson et al.¹ defined the criteria for success in implantology as follows: no mobility, pain, or peri-implant radiolucency. They also state in their definition that peripheral bone loss for the first year should be less than 1 mm and should subsequently be no more than 0.2 mm.

However, there are cases where further peripheral bone loss and peri-implantitis occur, but pathologic bone loss does not progress when an appropriate treatment is applied, and the implant can still handle a functional load. Such instances should not be defined as failures but rather as surviving im-

plants².

Survival and success rates have increased with the introduction of the rough-surface implant, which improves osseointegration³⁻⁵. In addition, with the use of screw-type and tapered implants, bone density has improved by intraosseous compaction; in this case, when biting force is applied, the stress is evenly dispersed into the peripheral bone tissues^{6,7}.

Rocci et al.⁸ reported that 2-3 years after machined surface implants were placed, 88% survived, whereas Lozada⁹ obtained a 95% success rate after an 8-year follow-up of hydroxyapatite-coated implants. Brechter et al.¹⁰ followed up patients for 30 months and obtained a survival rate of 98.5% for Ti-Unite (Nobel Biocare AG, Göteborg, Sweden) implants that had an anodizing oxidation surface. In contrast, Bornstein et al.¹¹ included a 6-week healing period after placement of an implant that had been sandblasted and given an acid-etched surface treatment, and obtained a 99.03% success rate at 3 years. According to a prospective study examining the stability of tapered resorbable blasting media (RBM) surface implants in the posterior maxilla, the 1-year survival rate was 97.4% and the success rate was 94.7%¹².

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However, the success and the survival rates of implants is determined not only by characteristics such as the structure and surface treatment of the implants but also by host factors such as age, gender, systemic disease, tobacco use or dental hygiene; biological factors such as bone disease, bone mass and bone grafting; and prosthetic factors such as the form of the prosthetic, occlusion and loading period¹³.

Romanos and Nentwig¹⁴ performed comparative studies of implants with immediate loading and implants with delayed loading, and reported survival rates of 94.9% and 91.68%, respectively. Becktor et al.¹⁵ found a significant difference in implant survival rate between patients who had a bone graft on the maxillary edentulous jaw and those who did not. They reported survival rates of 75.1% and 84.0%, respectively.

Shumaker et al.¹⁶ noted that if proper management, such as a regular clinical assessment, plaque control, and mouth hygiene, is not followed, inflammatory lesions such as peri-implantitis can occur despite aggressive treatment.

The purpose of this study was to evaluate the 7-year cumulative survival rate of the Osstem implants, US II, US III, SS II, and GS II (Osstem Implant Co., Ltd., Busan, Korea), with various structure and surface treatments in order to confirm their stability after placement and to analyze the factors that are involved in implant survival.

II. Materials and Methods

1. Subjects

The subjects of the study were patients who had Osstem implants that were placed at the Section of Dentistry, Seoul National University Bundang Hospital (Seongnam, Korea) from June 2003 through December 2005. One oral and maxillofacial surgeon and prosthodontist performed the treatment and we included patients who used tobacco as well as those with systemic disease. The study was conducted under Institutional Review Board approval of Seoul National University Bundang Hospital (B-1005-100-101). A total of 105 patients (55 male, 50 female) with a total of 467 Osstem implants were selected, and we examined their clinical records that contained historical charts and radiography. The observation period was between 42 and 83 months and began after placement of the final prosthesis.

2. Implant types and placement sites

The implants were divided into 4 types: SS II (straight

body, internal connection type, non-submerged type, RBM surface), US II (straight body, external connection type, RBM surface), US III (tapered body, external connection type, RBM surface), and GS II (straight body, internal connection submerged type and anodizing surface). Of these 4 types, 228, 199, 27, and 13 implants were respectively placed. For the US II implants, the implant of 4.0 mm in diameter was used most often, at 115 times; for the SS II implants, the 4.1 mm implant was used 122 times and the 4.8 mm implant was used 106 times. (Table 1) Implants of 11.5 mm in length were used most frequently in all product groups. (Table 2) In terms of positioning, 54 implants were placed in the anterior maxilla, 149 implants in the posterior maxilla, 65 implants in the anterior mandible and 199 implants in the posterior mandible. (Table 3)

3. Accompanying bone grafts and implant placement method

Accompanying bone grafts included a sinus bone graft, a ridge augmentation and a guided bone regeneration (GBR). There were 337 implants that were placed with bone grafts and 130 implants that were placed without bone grafts. In total, 158 implants were placed using the one-stage (non-submerged) method, and 309 implants were placed using the two-stage (submerged) method.

Table 1. Implant diameters

Diameter of fixture (mm)	Number
SS II	
4.1	122
4.8	106
Total	228
US II	
3.3	37
3.75	20
4.0	115
5.0	27
Total	199
US III	
4.0	14
5.0	13
Total	27
GS II	
3.5	3
4.0	5
4.5	3
5.0	2
Total	13

SS II, US II, US III, and GS II: Osstem implants (Osstem Implant Co., Ltd.).

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Table 2. Implant length

Length of fixture (mm)	Number
SS II	
8.5	10
10.0	27
11.5	119
13.0	61
15.0	11
Total	228
US II	
8.5	15
10.0	25
11.5	72
13.0	64
15.0	23
Total	199
US III	
10.0	2
11.5	17
13.0	8
Total	27
GS II	
10.0	2
11.5	8
13.0	1
15.0	2
Total	13

SS II, US II, US III, and GS II: Osstem implants (Osstem Implant Co., Ltd.).

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Table 3. Distribution of implant position

Position	Number
Maxilla	
Anterior	54
Posterior	149
Total	203
Mandible	
Anterior	65
Posterior	199
Total	264

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4. Prosthetic loading

In this study, we defined the conventional loading period for the lower jaw as 3 months and that for the upper jaw as 6 months; early loading and delayed loading were defined before and after the conventional loading period, respectively. The period before prosthesis loading ranged from 1 to 42 months and averaged 8.93 ± 7.35 months. For the upper jaw, the period before prosthesis loading ranged from 1 to 41 months, with an average of 10.35 ± 7.25 months, whereas these values for the lower jaw were 1 to 42.17 months, with an average of 7.81 ± 7.26 months.

5. Evaluation of prognosis

In determining prognosis for survival analysis, the following outcomes were regarded as failures: implant mobility, pain on percussion, presence of radiolucency around the implant on radiography and crestal bone loss of 1 mm or more over a period of one year². Periapical radiography was performed immediately after implantation, and these radiographs were compared with annual X-rays; we measured the crestal bone loss in the mesial and distal aspects and calculated the average.

6. Statistics

In this study, we used the life table method and a cross-tabulation analysis to analyze the cumulative survival rate of all 467 implants. We used the log rank test to evaluate the survival curve, and the influence that the prognostic factors, i.e., patient age, gender, bone graft history, implant diameter and width, loading time, implant type, and one-stage/two-stage implant, had on the survival rate was determined with a Cox proportional hazard model based on logistic regression analysis. Events were defined as cases in which implants were removed because of failure, and censored cases included those in which the implants survived until the observation concluded and those in which the patients were lost to follow-up or dropped out. We used PASW Statistics 18.0 for Windows (IBM Co., Armonk, NY, USA) as our statistics tool and the 95% confidence level to test the significance.

III. Results

1. Seven-year cumulative survival rate

Of all 467 implants, 20 implants were removed because of failure. Removal occurred as follows: 1 implant was removed 2 months after the surgery, 3 implants after 3 months, 4 implants after 4 months, 2 implants after 5 months, 1 implant after 6 months, 1 implant after 9 months, 1 implant after 13 months, 3 implants after 14 months, 2 implants after 15 months, and 2 implants after 29 months. The cumulative survival rate for each interval was 97.34% at 1 year, 95.90% at 2 years, and 95.37% at 3-7 years, and the final cumulative survival rate was 95.37%. (Table 4, Figs. 1, 2)

2. Comparison according to the type of implant

In groups US III and GS II, 100% of the implants survived.

Table 4. Seven-year cumulative survival analysis (total: 467, censored: 447, event: 20)

Interval time (yr)	Proportional surviving	Cumulative survival rate ¹ (%)	Hazard rate
1	0.9734	97.34	0.0022
2	0.9852	95.90	0.0012
3	0.9945	95.37	0.0005
4	1.0000	95.37	0.0000
5	1.0000	95.37	0.0000
6	1.0000	95.37	0.0000
7	1.0000	95.37	0.0000

¹Life table method.

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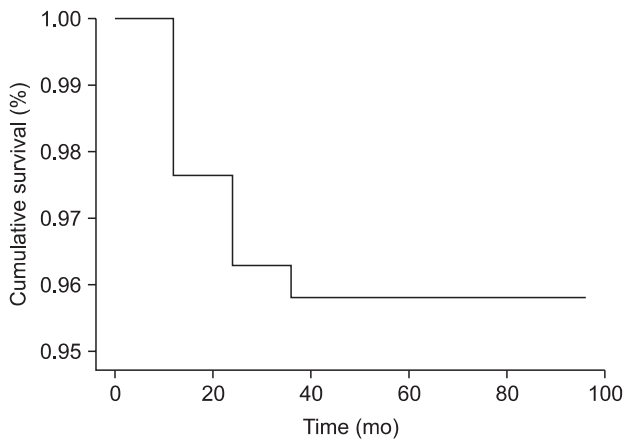


Fig. 1. Survival function curve.

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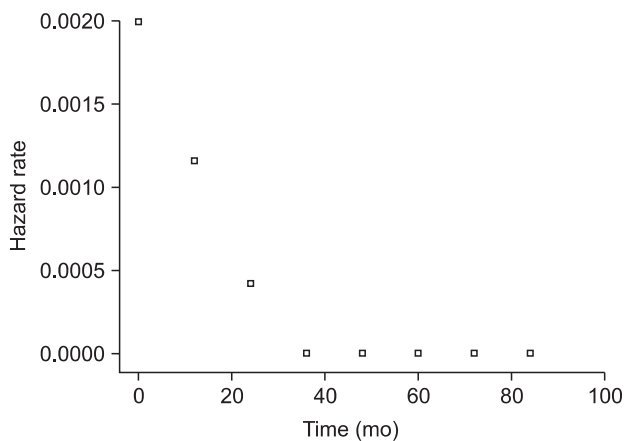


Fig. 2. Hazard function curve.

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In group SS II, the cumulative survival rate for each interval was 95.07% at 1 year and 93.62% at 2-7 years, and the final cumulative survival rate was 93.62%. In group US II, the

Table 5. Comparison of subgroups according to implant type

Interval time of subgroup (yr)	Cumulative survival rate ¹ (%)	Significance level ²
SS II		0.262
1	95.07	
2	93.62	
3	93.62	
4	93.62	
5	93.62	
6	93.62	
7	93.62	
US II		
1	99.48	
2	97.78	
3	96.55	
4	96.55	
5	96.55	
6	96.55	
7	96.55	
US III		
1-7	100	
GS II		
1-7	100	

SS II, US II, US III, and GS II: Osstem implants (Osstem Implant Co., Ltd.).

Quantities of implants: SS II: 228, US II: 199, US III: 27, GS II: 13.

¹Life table method. ²Log-rank test.

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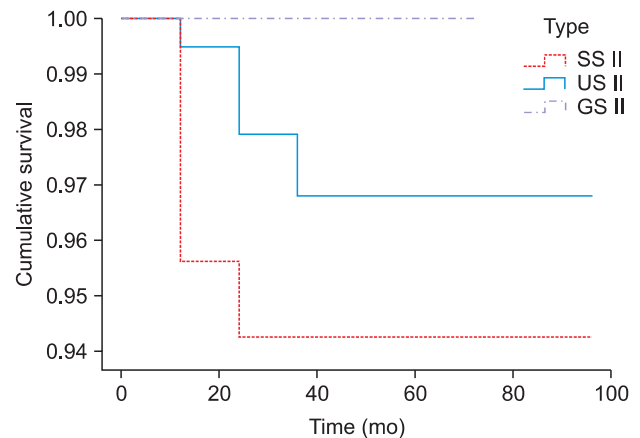


Fig. 3. Comparison of the survival function curves between subgroups. SS II, US II, and GS II are Osstem implants (Osstem Implant Co., Ltd.).

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cumulative survival rates for each interval were 99.48% at 1 year, 97.78% at 2 years and 96.55% at 3-7 years, and the final cumulative survival rate was 96.55%. In a comparison of the cumulative survival rate for each group using cross-tabulation analysis ($P>0.05$) and logistic regression analysis ($P>0.05$), we found that there was no statistically significant difference. (Table 5, Fig. 3)

3. Analysis of the prognostic factors that influence the cumulative survival rate

The cumulative survival rate was related to the diameter of the implant, the bone graft history and the loading period. Increased implant diameter, bone grafting, and a shorter healing time were factors that increased the risk of failure.(Table 6) In this study, among the 467 implants, the failure rate of implants placed in the maxilla was higher than for those placed in the mandible. In addition, 17 implants failed in cases that were accompanied by surgery with GBR. We applied early loading to 18 implants, regular loading to 210 implants and

delayed loading to 239 implants. Of the 20 failed implants, 7 implants had early loading, 10 implants had regular loading and 3 implants had delayed loading. Ten failed implants occurred in two months of loading in one patient. The failure rate of early-loaded implants was 50% in the mandible and 87.5% in the maxilla. Defining ‘wide-type’ implants as those with diameters ≥ 4.8 mm¹⁷, 10 of the 20 implant failures were wide-type implants and 10 implants were standard implants.(Table 7)

IV. Discussion

The survival of implants placed in an edentulous jaw is influenced by implant characteristics, biological environment of the host and prosthetic restoration; implant survival has improved as implant structure, surface treatment and bone graft techniques have improved. Cavallaro¹⁷ reported that when 176 implants were placed on a normal alveolar bone, the survival rate at 3 years was 98.6%, and when 28 implants were immediately placed in the wound of the tooth extraction, the survival rate was 96.4%. Stanford et al.¹⁸ placed 1,246 implants and analyzed the cumulative survival rate for a year after prosthetic loading (98.6%). Stafford¹⁹ revealed that when implantation occurred immediately after tooth extraction, the survival rate was 95.5%.

Table 6. Evaluation of other prognostic factors

Factor	Significance level ¹	Exp (B)	95% CI	
			Lower	Upper
Age	0.721	0.995	0.956	1.034
Sex	0.105	0.450	0.168	1.208
Diameter	0.021*	1.516	0.907	2.533
Length	0.963	1.134	0.797	1.613
Bone graft	0.046	0.494	0.195	1.247
1-Stage/2-Stage	0.547	1.466	0.524	4.103
Loading	0.000*	0.682	0.559	0.832

(CI: confidence interval)

¹Cox proportional hazard model.

*P<0.05.

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Table 7. Cases of implant failure

Age (yr)	Sex	ASA score	Site ¹	Type	Width (mm)	Loading (mo)	Surgery
72	F		3	SS II	4.1	2	
42	M		4	SS II	4.1	5	GBR
			4	SS II	4.1	2	GBR
45	F		2	US II	5.0	4	Sinus bone graft
70	F		2	SS II	4.1	13	
73	F		2	SS II	4.8	6	OSFE
			2	SS II	4.8	6	OSFE
54	M		2	SS II	4.1	8	Sinus bone graft
46	M		2	SS II	4.8	5	Sinus bone graft
29	F		4	SS II	4.8	13	
62	M	II (DM)	1	SS II	4.8	2	Le Fort I
			2	SS II	4.8	2	Iliac bone graft
			1	SS II	4.1	2	Delayed placement
			1	SS II	4.1	2	
			2	SS II	4.8	2	
			2	US II	4.0	2	
			2	US II	4.0	2	
			2	US II	5.0	5	
			2	US II	5.0	5	
			1	US II	5.0	5	

(ASA: American Society of Anesthesiologists, F: female, M: male, DM: diabetes mellitus, GBR: guided bone regeneration, OSFE: osteotome sinus floor elevation)

¹Site: 1 (anterior maxilla), 2 (posterior maxilla), 3 (anterior mandible), 4 (posterior mandible).

SS II, US II: Osstem implants (Osstem Implant Co., Ltd.).

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Survival analysis is a statistical technique in which the survival period is analyzed using a survival function, and the distribution of the survival period of a group, including members whose observation has not been completed, can be determined. In this study, the cumulative survival rates of Osstem implants were 97.34% at 1 year after placement, 95.90% after 2 years, and 95.37% after 3 years, which were not largely different from previous studies. According to a multicenter clinical study of Osstem implants after maxillary sinus floor elevation, the survival rates of US II and SS II implants were 99.2% and 95.8%²⁰. A multicenter clinical study of Osstem GS II implants reported that the success and survival rates of the implants were 93.23% and 95.83%, respectively, in type IV bone²¹.

Failure of the implant in the early stage is related to the primary stability of the implant, which can be improved by selecting implants with appropriate structure and surface properties as well as by improving the surgical method²². Osstem implants include implants with various structures depending on the gingival biotype, bone quality and bone defects; in the US III and GS II groups, 100% of implants survived; in the US II group, the final cumulative survival rate was 96.55%; and in the SS II group, the final cumulative survival rate was 93.62%.

Statistical analysis for the cumulative survival rate was performed using cross-tabulation ($P > 0.05$) and logistic regression ($P > 0.05$) analyses, but there were no statistically significant differences between any of the groups. However, because the numbers of followed US III and GS II implants were substantially smaller than those of the SS II and US II implants and because 100% of the US III and GS II implants survived, a comparative analysis on the differences between the SS II and US II implants is required.

The cross-tabulation and logistic regression analyses revealed no statistically significant differences between the SS II and US II implants. The US II implant has an external hex connection structure and is of the submerged type; because of the RBM surface treatment to improve the affinity with bone, this implant can easily be placed in various bone qualities and has outstanding holding power. In contrast, the SS II implant has an internal 8 degree morse taper and is of the non-submerged type; it has a stable connection structure and can be easily placed in various bone qualities. The GS II implant has an internal hex connection structure and is of the submerged type; its advantage is a dual micro and macro thread that minimizes bone resorption and provides optimal stress dispersion. The US III implant has an RBM surface

and is of the tapered type with an external connection structure. When placed on a site with weak bone quality, powerful bone compression can be obtained that is effective at 1/3 of the retainer^{23,24}.

Of the many factors that may influence the survival of implants, we confirmed that patient age and gender as well as the length of the implants, choice of one-stage method or two-stage method, and the type of product were not related to increased risk. Although it has been suggested that failure rate increases as age increases because of decreased bone density caused by bone resorption exceeding osteogenesis²⁵, there was no increased risk related to age in this study as well as in the study of Smith et al.²⁶. There was no difference in risk between males and females, as previously determined by Heberer et al.²⁷. Renouard and Nisand²⁸ reported that, by avoiding countersink or under-preparation at sites where the bone substance was poor, there was no difference in the success rate between rough-surface implants that were greater than 10 mm and less than 10 mm. In the present study, we also did not identify any risk related to implant length.

Some scholars have claimed that having a bone graft history was not a risk factor for implant failure^{29,30}. However, a more serious surgical wound is more likely to affect the peripheral blood supply and therefore have a harmful influence on the healing of soft tissue and bone tissue. Becktor et al.¹⁵ found that the implant survival rates were 75.1% and 84.0% depending on whether or not there was a bone graft on the maxillary edentulous jaw, respectively, which represented a significant difference. Some authors have suggested that an implant placed in an adequate recipient site should perform better than an implant placed in a reconstructed site³¹. In this study, a history of bone grafting was directly correlated with increased risk, and we suggest that surgical techniques should focus not only on survival, but also on procedures that are less complicated, less invasive and are accompanied by smaller complications.

In addition, risk of failure increased with increasing implant diameter and earlier loading period. Ivanoff et al.³² reported an increased failure rate for implants with a wide diameter (≥ 5 mm) and attributed this effect not only to the difference in surgical technique, but also to the inability to obtain primary stability due to poor bone quality, which resulted in the use of wide implants as 'rescue' implants. Wide-diameter implants are often used when the bone substance and bone mass are poor, which increases the early failure rate.

Optimal loading time for upper dental prostheses after implant placement is still debated. The 2002 World Con-

gress Consensus Meeting in Barcelona defined immediate loading as 3-4 days after surgery³³. In addition, the 2003 ITI Consensus Conference defined the period within first 2 days after placement as immediate loading, the period between 48 hours and 3 months as early loading and the period between 3 months and 6 months as conventional loading³⁴. At the 2008 ITI conference³⁵, immediate loading referred to less than 1 week after placement, early loading referred to between 1 week and 2 months after placement and conventional loading referred to 2 months or more after placement. Under the same conditions, many other clinical studies as well as histomorphometric observations have confirmed better prognosis for delayed loading than for early loading³⁰. In this study, similar to past findings, we observed that the risk of failure decreased as the loading time was delayed.

Of the 20 cases in which the implants were removed due to failure, 1 case was attributed to a psychiatric cause, and the remaining 19 were caused by the failure of osseointegration. Out of the 20 failed implants, 50% occurred in 1 patient. This patient had controlled DM as well as severe maxillary alveolar bony deficiency; therefore, he underwent interpositional bone grafting using iliac bone, and delayed implantation took place 4 months following the bone graft. Extensive bone grafting and surgical wounds are considered to be major factors in implant failure; the concentration of implant failures in one patient has been called 'clusterization' in previous studies^{36,37}.

We suggest that the cause for this clusterization likely involved an immunological reaction of the patient in response to the systemic disease, the overloading of the bone substance or insufficient bone mass due to oral parafunction such as bruxism, the use of a difficult surgical technique or the application of early loading. In addition, of the 20 failed implants, 14 implants were SS II and 6 implants were US II; of the 6 US II implants, 5 implants occurred in the aforementioned patient.

One limitation of this study is that the factors that influenced implant survival were broad because we chose a retrospective study design. This approach limited our scope of research, and we did not consider the prosthetic aspect. In this study, we only considered the patient's age, gender, bone graft history, implant diameter and width, product type, one-stage/two-stage implant, and loading time as factors that influenced survival of implants. In addition to these factors, other factors such as the bone graft material, membrane, systemic disease, surgery time, type of anesthesia, type of prosthesis, condition of the occluding dentition, parafunction,

and tobacco use also influence the survival rate, but were excluded in this study.

If we consider that each variable is interactional, then a duplicate statistical model, such as multi-regression analysis, would be required. However, because of the characteristics of the data, which have more discrete variables than continuous variables, we determined that it would be difficult to obtain a satisfactory result with this method. Therefore, we used cross-tabulation analysis and logistic regression analysis instead, which are widely used. A prospective study in the future will eliminate factors that influence the cumulative survival rate of implants and their prognosis while examining each factor independently.

V. Conclusion

The seven-year cumulative survival rate of Osstem implants was 95.37%. As the healing period after implant placement increased, the failure risk decreased. However, as the diameter of the implants increased, the failure risk also increased, depending on bone graft history and early loading.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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