

Long-term Survival and Success of Oral Implants in the Treatment of Full and Partial Arches: A 7-year Prospective Study with the ITI Dental Implant System

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Purpose: This study evaluated the long-term survival and success of different implant-supported prostheses supported by ITI implants. **Materials and Methods:** Two hundred fifty consecutive patients were rehabilitated using implant-supported prostheses. Seven hundred fifty-nine implants were loaded. Single-tooth prostheses ($n = 106$), cantilever fixed partial prostheses ($n = 42$), fixed partial prostheses ($n = 137$), fixed complete prostheses ($n = 5$), implant/tooth-supported prostheses ($n = 13$), and overdentures ($n = 37$) were used. The mean follow-up period was 3.85 years. Life table analyses were performed. Implant survival rates were calculated by means of standard life table principles. Statistical analysis was performed to compare the implant survival and success by implant placement site for each type of prosthesis. **Results:** The cumulative implant survival rates were calculated for implants supporting single-tooth prostheses (95.6%), cantilever fixed partial prostheses (94.4%), fixed partial prostheses (96.1%), fixed complete prostheses (100%), implant/tooth-connected prostheses (90.6%), and overdentures (95.7%). Similar survival and success rates were documented for implants placed in maxillae and mandibles. Implant size did not influence survival. **Discussion:** Seven-year survival rates were similar for implants supporting single-tooth prostheses, cantilever fixed partial prostheses, fixed partial prostheses, and implant/tooth-supported prostheses. Medium-long term implant survival and success were not influenced by the site (maxilla or mandible). Implant and prosthetic survival rates for overdentures supported by 2 implants were comparable to those for overdentures supported by 3 or more implants. **Conclusion:** Prostheses supported by ITI implants represent a reliable medium-term treatment. (More than 50 references.) INT J ORAL MAXILLOFAC IMPLANTS 2004;19:247-259

Key words: dental implants, implant-supported restorations, peri-implant bone resorption

Implant dentistry is a valid and predictable treatment option for the rehabilitation of partially and completely edentulous arches. An important contri-

bution to this field has been provided by the continuous quality improvement of prosthetic components. At present, a number of implant-supported prosthetic solutions can satisfy patients' expectations regarding esthetics and function. The success of the ITI implant system (Straumann, Waldenburg, Switzerland) is well documented in international literature, particularly regarding the treatment of completely edentulous arches.¹⁻¹³ Prospective and longitudinal studies related to partial edentulism indicate cumulative implant success rates ranging from 89% to 95.3%, and cumulative survival rates ranging from 93.6% to 96.7%, 3 to 7 years after loading.¹⁴⁻¹⁷

The purpose of this study was to evaluate the medium to long-term survival and success of different implant-supported prostheses supported by ITI implants. Another purpose was to determine whether significant differences in survival and success could be observed for different implant placement sites.

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Table 1 Implant Placement Location

Location	n	%
Maxillary		
Anterior	112	14.8
Posterior	223	29.4
Mandibular		
Anterior	99	13.0
Posterior	325	42.8
Total	759	100.0

*Two mandibular posterior implants were removed before prosthetic loading. Seven hundred fifty-seven implants were loaded and followed up.

Table 2 Implant Features

Parameter	n
Height	
8 mm	72
10 mm	402
12 mm	236
14 mm	48
16 mm	1
Diameter	
3.3 mm	149
4.1/4.0 mm	579
4.8 mm	31
Type	
Hollow screw	195
Solid screw	564
Surface	
Titanium plasma-sprayed	703
Sandblasted acid-etched	56

MATERIALS AND METHODS

Patient Selection

Two hundred fifty patients (106 men and 144 women between 20 and 67 years of age) were consecutively treated between May 1995 and June 2002. Patients were treated at the Dental Clinic, Department of Medicine Surgery and Medicine, University of Milan, Italy. All patients signed an informed consent form. After prosthesis placement, the patients were seen for follow-up for periods ranging from 16 months to 7 years (mean 3.85 years).

Inclusion criteria were:

- Partial or complete edentulism
- Adequate bone volume at implant site for an implant at least 3.3 mm in diameter and 8 mm in length. Bone volume was evaluated by juxtagin-gival radiographs and clinical judgment.
- A favorable maxillomandibular relationship

Exclusion criteria were:

- Periodontal disease
- Heart disease
- Coagulation or leukocytic diseases
- Metabolic disorders
- Radiotherapy in the head or neck area
- Parafunctions such as clenching or bruxism
- A smoking habit (more than 10 cigarettes per day) or alcohol or drug abuse¹³

Implants and Prostheses

Patients were restored with ITI implant-supported prostheses. Seven hundred fifty-nine implants were placed. Two implants were removed before prosthetic loading. During the 7-year follow-up period, 49 patients with 82 implants and 45 restorations were lost to follow-up. Five patients died after they had received fixed prostheses; 4 patients developed severe illness that prevented further follow-up; 24 patients moved and dropped out of the study; and 14 patients could not be reached. Two patients were transferred to a private practice but never attended scheduled follow-up visits. The corresponding implants have been classified as “dropouts” and are unaccounted for.

The distribution of the implants is shown in Table 1. Three hundred thirty-five implants were placed in the maxilla and 424 were placed in the mandible. Loaded implants ranged from 8 to 16 mm in length and from 3.3 to 4.8 mm in width (Table 2).

One hundred thirty-seven prostheses were placed in the maxilla and 203 were placed in the mandible. The following types were used (Table 3): 110 fixed single-tooth (ST) prostheses (Fig 1a), 42 cantilever fixed partial (CFP) prostheses (24 with mesial cantilever and 18 with distal) (Figs 1b and 1c), 137 fixed partial (FP) prostheses (Fig 1d), 5 fixed complete (FC) prostheses (Figs 1e and 1f), 13 implant/tooth-supported (ITS) prostheses (Fig 1g), and 37 overdentures (ODs) (Figs 1h and 1i).

Clinical and Radiographic Assessment

Peri-implant tissue was evaluated for suppuration,¹⁹ probing pocket depth and probing attachment level,²⁰ bleeding on probing (Bleeding Index score of 0 to 3),²¹ and peri-implant inflammation (using the guidelines suggested by Mombelli and Lang).²² Plaque Index scores (on a scale of 0 to 3),²¹ implant mobility (either given a score of 0 to 2 [see Mombelli and coworkers²¹] or evaluated using the Periotest [Siemens, Bansheim, Germany]), and percussion¹⁸ were also noted.

Intraoral radiographs were taken using the paralleling technique to control projection geometry,

Table 3a Implants			
	Maxilla	Mandible	Total
Prosthesis supported			
Single tooth	44	62	106
Cantilever fixed partial	40	44	84
Fixed partial	121	174	295
Fixed complete	21	14	35
Implant/tooth-supported	22	9	31
Overdenture	40	74	114
Total	288*	377*	665*

*Implants of patients considered to be dropouts are excluded.

Table 3b Prostheses			
	Maxilla	Mandible	Total
Prosthesis			
Single tooth	44	62	106
Cantilever fixed partial	20	22	42
Fixed partial	53	84	137
Fixed complete	3	2	5
Implant/tooth-supported	8	5	13
Overdenture	9	28	37
Total	137*	203*	340*

*Implants of patients considered to be dropouts are excluded.



Fig 1a Juxtagingival radiograph of an ST prosthesis.



Fig 1b Juxtagingival radiograph of a fixed prosthesis cantilevered mesially.



Fig 1c Juxtagingival radiograph of a fixed prosthesis cantilevered distally.

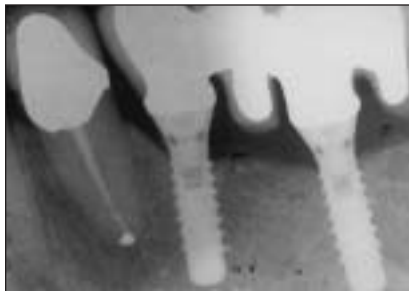


Fig 1d Juxtagingival radiograph of an FP prosthesis.



Fig 1e Cover screws used during healing for implants supporting an FC prosthesis.



Fig 1f Loading of an FC prosthesis.



Fig 1g Juxtagingival radiograph of an ITS prosthesis.



Fig 1h Bar anchorage of 4 implants used for OD support.



Fig 1i Ball attachment anchorage of 2 implants used for OD support.

with exposure parameters of 65 to 90 kV, 7.5 to 10 mA, and 0.22 to 0.25 seconds.²³⁻²⁹ Radiographs were taken at loading, at 6 and 12 months after loading, and annually thereafter. They were subsequently stored on a personal computer and analyzed using the program ImageJ (US National Institute of Mental Health, Bethesda, MD) to determine bone level changes. Peri-implant bone resorption was assessed at bone levels mesial and distal to the implants.

Prosthetic Treatment

Patients with type 1 or type 2 bone quality were recalled for a preprosthetic evaluation 3 months after implant placement; patients with type 3 or type 4 bone quality were recalled after 6 months.³⁰ After the removal of the cover screws used during healing 3 to 6 months after implant placement, the abutments were screwed into position using 35 Ncm of torque. Solid abutments or Octa abutments (Straumann) with cast-to parallelized gold copings were used for cemented prostheses. Octa abutments with cast-to gold copings were used for screw-retained prostheses.

For ODs, the framework and esthetic veneer were fabricated from gold alloy and resin; all other prosthetics were fabricated from gold alloy and porcelain. No soldering was performed. Cemented prostheses were fixed with zinc oxyphosphate cement or zinc-eugenol oxide cement, while screw-retained prostheses were secured by applying a 15-Ncm torque to the abutment-framework screw using a manual torque driver.

Statistical Analysis

The statistical analysis was performed with the life table analysis described by Kalbleish and Prentice³¹ and Colton.³² The data were analyzed at the end of December 2002. Thus, all restored implants completed at least the 1-year examination. The following analyses were performed for the entire group of 759 implants:

- *Cumulative survival rate:* The internal survival rate for each time interval and the cumulative survival rate for the entire 8-year period were calculated. In the present study, survival rate was defined as the percentage of load-bearing implants that did not show symptoms of pain, mobility,²¹ or infection (not including successfully treated peri-implant inflammation).²²
- *Cumulative success rate:* In this analysis, which was more strict than the survival rate analysis, implants exhibiting a suppurative peri-implant infection at the last annual examination were classified as “failures.” In addition to failed

implants, this index also took into account the failure criteria established by Albrektsson and colleagues,³³ Buser and colleagues,³⁴ and van Steenberghe.³⁵

Life tables included the following parameters: time period (observation time), number of implants at interval start (O^x), number of early failures (ie, implants that failed prior to loading), number of loaded implants, number of implants lost to follow-up as a result of patient dropout (w^x), number of implants at risk (ie, the “harmonic mean” of the implants at the beginning of an interval and those remaining at the end), number of failed implants (d^x) in each interval (q^x), interval survival rate, and cumulative survival rate.³² The interval failure rate formula was

$$q^x = \frac{d^x}{O^x - \frac{w^x}{2}}$$

The interval survival rate (p^x) formula was $p^x = 1 - q^x$. However, the authors have accounted for the correlation structure associated with multiple implants in the same subject. Thus, unless the patient was restored with a single implant supporting a single crown, the reduced significance of any comparison was considered.

A 4-field table^{36,37} was used to present the number of successes, survivals, unaccounted-for implants, and failures up to the 7-year examinations. The matrix gave a qualitative description of the materials and the methods of evaluation used and presented the results. The failure rate was always cumulative, and the sum of the 4 fields was 100% (ie, all placed implants).

Prognostic Criteria

The following criteria for implant evaluation were considered³⁷:

- *Failure:* An implant was regarded as a failure if it had to be removed for any reason. If there was clinical mobility, implant removal was indicated without question. Absence of mobility was checked by applying a light tightening force to the implant with an abutment screwdriver without simultaneously counteracting the force with an abutment clamp. Any mobility or sensation (eg, pain) was regarded as a sign of lost osseointegration. Other conditions for which implant removal could be indicated included, for example, incurable soft tissue infection, persistent pain, paresthesia, or discomfort. Information

regarding any adverse event, including condition on onset, measures taken, and recovery, was recorded. Adverse events did not always result in removal.

- *Survival*: Implants that did not fail were included in the survival group.
- *Success*: Surviving implants were considered successful if radiographic evaluation revealed no more than 1.0 mm of marginal bone loss during the first year of loading and no more than 0.2 mm resorption per year in subsequent years, if there was no peri-implant pathosis or radiolucency, and if probing depth (measured with a calibrated plastic probe) was no greater than 3 mm on any side (mesial, buccal, distal, or lingual-palatal). Juxtagingival radiographs taken at the time of abutment connection and at the time of the last follow-up were used for measurements. All successful implants were in clinical function.

RESULTS

Results of the 7-year Life Table Analysis

Two mandibular implants were lost before loading because of persistent paresthesia and were classified as “early failures.” During the 7-year follow-up period, 30 implants failed (Table 4). Implant mobility was recorded for 27 implants (14 maxillary, 13 mandibular) in 21 patients (Figs 2 and 3) owing to severe peri-implantitis (n = 21) and biomechanical overloading (n = 6). Three maxillary implant fractures occurred—two in implants supporting CFP prostheses and 1 in an implant supporting an FP prosthesis. All failed implants were removed. Six failed implants were short (8 mm in height) and 8 were narrow (3.3 mm in diameter); 1 failed implant was short and narrow.

Twenty-eight cases of peri-implantitis were successfully treated.²² Thirty-five implants showed more than 1.0 mm of marginal bone loss during the first year of loading, followed by more than 0.2 mm bone resorption per year. Seven implants had a peri-implant probing depth greater than 3 mm.

Cumulative survival rates were calculated for implants supporting ST prostheses (95.6%), CFP prostheses (94.4%), FP prostheses (96.1%), FC prostheses (100%), ITS prostheses (90.6%), and ODs (95.7%) as shown in Table 5.

The 4-field analysis included calculation of the 7-year failure rate for implants supporting ST prostheses (4.0%), CFP prostheses (5.4%), FP prostheses (3.6%), FC prostheses (0%), ITS prostheses (8.8%), and ODs (4.0%), as shown in Table 6. Dropout rates were 13.8% for implants supporting ST prostheses,

Table 4 Failed Implants

Reason failed	Diameter (mm)	Height (mm)	Prosthesis supported	Location
Implant fracture (biomechanical overloading)				
1	3.3	10	CFP	Maxilla
2	3.3	10	CFP	Maxilla
3	4.1	12	FP	Maxilla
Implant mobility (biomechanical overloading)				
1	4.1	12	OD	Mandible
2	4.1	10	OD	Mandible
3	4.1	8	ST	Mandible
Implant mobility (peri-implantitis)				
1	4.1	10	FP	Mandible
2	3.3	10	FP	Mandible
3	4.8	10	FP	Mandible
4	4.1	10	FP	Mandible
5	3.3	8	FP	Mandible
6	4.1	10	FP	Maxilla
7	4.1	8	FP	Maxilla
8	3.3	10	FP	Maxilla
9	4.1	10	FP	Maxilla
10	4.1	8	FP	Maxilla
11	4.1	10	FP	Maxilla
12	4.8	10	CFP	Mandible
13	4.1	12	CFP	Mandible
14	3.3	10	CFP	Maxilla
15	4.1	10	ST	Maxilla
16	4.1	8	ST	Maxilla
17	4.1	12	ST	Mandible
18	3.3	10	ST	Mandible
19	4.1	10	OD	Maxilla
20	4.1	10	OD	Maxilla
21	4.1	10	OD	Maxilla
22	4.1	12	ITS	Mandible
23	4.1	8	ITS	Maxilla
24	3.3	10	ITS	Maxilla
Persistent paresthesia				
1	4.1	10	Not loaded	Mandible
2	4.1	10	Not loaded	Mandible

9.7% for CFP prostheses, 12.2% for FP prostheses, 25.5% for FC prostheses, 8.8% for ITS prostheses, and 9.5% for ODs. The proportion of successes after 7 years was 75.6% for implants supporting ST prostheses, 76.3% for CFP prostheses, 73.8% for FP prostheses, 63.8% for FC prostheses, 70.6% for ITS prostheses, and 78.6% for OD prostheses.

Prosthetic complications such as decementation (4 ST and 5 FP prostheses), abutment-framework connection screw loosening (3 FP prostheses), retention anchor failure (n = 3), and retaining screw loosening (2 OD prostheses) also occurred. One abutment screw and 1 prosthetic pontic, both in FP prostheses, fractured; 6 aesthetic veneer fractures (2 in ST prostheses and 4 in FP prostheses) were observed. OD fracture was recorded twice in 2 different patients (Table 7).



Fig 2 Juxtagingival radiograph of a failed implant. The failure was related to biomechanical overloading.



Fig 3 An implant whose failure was related to peri-implantitis. Note the peri-implant bone resorption and the exposure of the implants threads.

Maxillary and mandibular implants showed similar outcomes. Mandibular implants demonstrated 7-year cumulative survival rates of 95.6% for ST prostheses, 95.7% for CFP prostheses, 97.2% for FP prostheses, 100% for FC prostheses, 89.9% for ITS prostheses, and 97.5% for OD prostheses (Table 8). Maxillary implants (Table 9) showed cumulative survival rates of 95.7% for ST prostheses, 92.8% for CFP prostheses, 95.6% for FP prostheses, 100% for FC prostheses, 91.7% for ITS prostheses, and 92.5% for ODs.

Finally, the outcomes of implants supporting ODs supported by 2 implants were compared to those of implants supporting ODs supported by 3 or more implants. The former group had a cumulative survival rate of 93.6%; the latter had a cumulative survival rate of 96.7% (Table 10).

DISCUSSION

Overall, implant and prosthesis success and survival rates were satisfactory; however, these data had a relative value. Prosthetic restorations were in fact exposed to qualitatively and quantitatively different complications. Literature studies^{30,31} have demonstrated higher survival probability for osseointe-

grated mandibular implants than for maxillary implants. The reasons for this are likely the lower mechanical stress that the maxilla can withstand because of its thinner cortical layer, as well as the lower density of the maxillary spongiosa.³⁷

Further, Scurria and associates³⁸ reported outcomes of dental implant therapy for a range of patients requiring various degrees of tooth replacement. Life table analyses produced Kaplan-Meier survival curves for different locations and prosthesis types. Proportional hazards modeling combined with SUDAAN modeling (RTI International, Research Triangle Park, NC) identified removable prosthesis type and maxillary location as being significantly associated with implant failure. For this reason, the present study analyzed implant survival and success by prosthesis type and implant location (maxilla or mandible).

Study results were comparable to those found in the international literature related to ST prostheses,^{13,15,17,38} FP prostheses,^{17,39-42} CFP prostheses,⁴³ ITS prostheses,^{44,45} FC prostheses,^{42,46,47} and ODs.^{4,8,48-50}

It was also noted that implant location did not represent a significant prognostic factor; the works of Adell and coworkers,⁵¹ Bass and Triplett,⁵² Jaffin and Berman,⁵³ and Jemt and coworkers⁵⁴ support

Table 5 Life Table Analysis for Implant Survival

Interval (y)	Implants at start of interval	Early failures	Loaded implants	Dropouts	Implants at risk	Failures during interval	Survival rate (%)	Cumulative survival rate (%)
ST prostheses								
0 to 1	123	2	121	3	119.5	0	100	100
1 to 2	118	0	118	0	118	0	100	100
2 to 3	118	0	118	5	115.5	3	97.4	97.4
3 to 4	110	0	110	3	108.5	2	98.2	95.6
4 to 5	105	0	105	2	104	0	100	95.6
5 to 6	103	0	103	0	103	0	100	95.6
6 to 7	103	0	103	2	102	0	100	95.6
CFP prostheses								
0 to 1	93	0	93	0	93	0	100	100
1 to 2	93	0	93	0	93	0	100	100
2 to 3	93	0	93	3	91.5	1	99	99
3 to 4	89	0	89	2	88	0	100	99
4 to 5	87	0	87	4	85	2	97.7	96.7
5 to 6	81	0	81	0	81	2	97.6	94.4
6 to 7	79	0	79	0	79	0	100	94.4
FP prostheses								
0 to 1	336	0	336	4	334	0	100	100
1 to 2	332	0	332	10	327	0	100	100
2 to 3	322	0	322	6	319	2	99.4	99.4
3 to 4	314	0	314	7	310.5	4	98.7	98.1
4 to 5	303	0	303	4	301	4	98.7	96.8
5 to 6	295	0	295	8	291	2	99.3	96.1
6 to 7	285	0	285	2	284	0	100	96.1
FC prostheses								
0 to 1	47	0	47	0	47	0	100	100
1 to 2	47	0	47	0	47	0	100	100
2 to 3	47	0	47	6	44	0	100	100
3 to 4	41	0	41	6	38	0	100	100
4 to 5	35	0	35	0	35	0	100	100
5 to 6	35	0	35	0	35	0	100	100
6 to 7	35	0	35	0	35	0	100	100
ITS prostheses								
0 to 1	34	0	34	0	34	0	100	100
1 to 2	34	0	34	0	34	0	100	100
2 to 3	34	0	34	1	33.5	0	100	100
3 to 4	33	0	33	0	33	1	97	97
4 to 5	32	0	32	1	31.5	1	96.8	93.8
5 to 6	30	0	30	1	29.5	0	100	93.8
6 to 7	29	0	29	0	29	1	96.6	90.6
OD prostheses								
0 to 1	126	0	126	0	126	0	100	100
1 to 2	126	0	126	2	125	0	100	100
2 to 3	124	0	124	2	123	0	100	100
3 to 4	122	0	122	2	121	2	98.4	98.4
4 to 5	118	0	118	4	116	0	100	98.4
5 to 6	114	0	114	2	113	0	100	98.4
6 to 7	112	0	112	0	112	3	97.3	95.7

Table 6 Implant Distribution After the 7-Year Follow-up

Prosthesis	Success n (%)	Dropout n (%)	Survival n (%)	Failure n (%)
ST	93 (75.6)	17 (13.8)	8 (6.6)	5 (4.0)
CFP	71 (76.3)	9 (9.7)	8 (8.6)	5 (5.4)
FP	248 (73.8)	41 (12.2)	35 (10.4)	12 (3.6)
FC	30 (63.8)	12 (25.5)	5 (10.7)	0 (0.0)
ITS	24 (70.6)	3 (8.8)	4 (11.8)	3 (8.8)
ODs	99 (78.6)	12 (9.5)	10 (7.8)	5 (4.0)

Table 7 Complications and Failures in Overdentures

	Anchors*		Bars†		Lock-pins‡	
	n (%)	Cause	n (%)	Cause	n (%)	Cause
Implant						
Complication	1 (3.1)	TP	5 (9.2)	PBR	4 (11.1)	TP
Failure	2 (6.2)	FIO	3 (5.5)	FIP	0	
Prosthetic						
Complication	3 (20.0)	RAR	2 (14.2)	RSL	0	
Failure	1 (6.6)	ODF	1 (7.1)	ODF	0	

*32 implants, 15 prostheses.

†54 implants, 14 prostheses.

‡36 implants, 8 prostheses.

PBR = peri-implant bone resorption exceeding success parameters; FIO = implant failure due to biomechanical overload; FIP = implant failure due to periimplantitis; TP = successfully treated peri-implantitis; RAR = retention anchors replacement; RSL = retaining screw loosening; ODF = overdenture fracture.

these findings. Nevertheless, in experimental and clinical literature, there are reports of implant survival and success being significantly influenced by placement site.^{38,55,56} Scurria and colleagues,³⁸ Romeo and colleagues,⁵⁵ and Balshi and colleagues⁵⁶ emphasized the greater biomechanical loads supported by posterior segments.

Comparing FP prostheses and CFP prostheses, statistically different implant and survival rates were not documented. This supports the reliability of a suitably fabricated cantilever prosthesis if the occlusal and biomechanical forces are appropriately addressed.

No notable difference was observed between survival rates of implant-supported FC prostheses and ODs, although the materials used to fabricate the suprastructure of an OD are usually more fragile than those used for an FC prosthesis, and ODs usually have a higher crown-to-implant ratio. The percentage of prosthetic complications was greater for ODs than for any other type of prosthesis (13.5% for ODs; 0% to 7.6% for other prostheses). This is probably

related to the relative weakness of the anchorage components used for connecting the implants to the prosthetic framework. The results of a number of studies on the functionality of ODs supported by 2 or more implants^{12,57} were partly confirmed. Implants supporting ODs supported by 2 implants survived nearly as well as implants supporting ODs supported by 3 or more implants.

Finally, implant failure did not appear to be significantly influenced by length and diameter; only 20% of failed implants were 8 mm long, and only 26.6% were 3.3 mm wide. Friberg and colleagues⁵⁸ published a long-term follow-up study of 49 patients in whom 260 short implants were placed. Their outcome showed the high predictability of that treatment procedure. Teixeira and coworkers⁵⁹ investigated the applicability of short hydroxylapatite-coated dental implants in the posterior mandible of partially edentulous patients. In a 5-year survival study, the overall cumulative implant survival rate was 94%, and the overall cumulative prosthetic survival rate was 91%. Predictable success

Table 8 Life Table Analysis for Survival of Loaded Mandibular Implants

Interval (y)	Implants at start of interval	Early failures	Loaded implants	Dropouts	Implants at risk	Failures during interval	Survival rate (%)	Cumulative survival rate (%)
ST prostheses								
0 to 1	72	2	70	2	69	0	100	100
1 to 2	68	0	68	0	68	0	100	100
2 to 3	68	0	68	2	68	1	98.5	98.5
3 to 4	64	0	64	1	63.5	2	97.1	95.6
4 to 5	62	0	62	2	61	0	100	95.6
5 to 6	60	0	60	0	60	0	100	95.6
6 to 7	60	0	60	1	59.5	0	100	95.6
CFP prostheses								
0 to 1	49	0	49	0	49	0	100	100
1 to 2	49	0	49	0	49	0	100	100
2 to 3	49	0	49	3	49	1	98	98
3 to 4	45	0	45	0	45	0	100	98
4 to 5	45	0	45	2	44	0	100	98
5 to 6	43	0	43	0	43	1	97.7	95.7
6 to 7	42	0	42	0	42	0	100	95.7
FP prostheses								
0 to 1	195	0	195	2	194	0	100	100
1 to 2	193	0	193	7	189.5	0	100	100
2 to 3	186	0	186	4	184	0	100	100
3 to 4	182	0	182	4	180	1	99.4	99.4
4 to 5	177	0	177	0	177	2	98.9	98.3
5 to 6	175	0	175	4	173	2	98.9	97.2
6 to 7	169	0	169	0	169	0	100	97.2
FC prostheses								
0 to 1	14	0	14	0	14	0	100	100
1 to 2	14	0	14	0	14	0	100	100
2 to 3	14	0	14	0	14	0	100	100
3 to 4	14	0	14	0	14	0	100	100
4 to 5	14	0	14	0	14	0	100	100
5 to 6	14	0	14	0	14	0	100	100
6 to 7	14	0	14	0	14	0	100	100
ITS prostheses								
0 to 1	10	0	10	0	10	0	100	100
1 to 2	10	0	10	0	10	0	100	100
2 to 3	10	0	10	1	9.5	0	100	100
3 to 4	9	0	9	0	9	0	100	100
4 to 5	9	0	9	0	9	0	100	100
5 to 6	9	0	9	0	9	0	100	100
6 to 7	9	0	9	0	9	1	89.9	89.9
OD prostheses								
0 to 1	84	0	84	0	84	0	100	100
1 to 2	84	0	84	2	83	0	100	100
2 to 3	82	0	82	0	82	0	100	100
3 to 4	82	0	82	2	81	2	97.5	97.5
4 to 5	78	0	78	4	76	0	100	97.5
5 to 6	74	0	74	2	73	0	100	97.5
6 to 7	72	0	72	0	72	0	100	97.5

Table 9 Life Table Analysis for Survival of Loaded Maxillary Implants

Interval (y)	Implants at start of interval	Early failures	Loaded implants	Dropouts	Implants at risk	Failures during interval	Survival rate (%)	Cumulative survival rate (%)
ST prostheses								
0 to 1	51	0	51	1	50.5	0	100	100
1 to 2	50	0	50	0	50	0	100	100
2 to 3	50	0	50	3	48.0	1	97.9	97.9
3 to 4	46	0	46	2	45	1	97.8	95.7
4 to 5	43	0	43	0	43	0	100	95.7
5 to 6	43	0	43	0	43	0	100	95.7
6 to 7	43	0	43	1	42.5	0	100	95.7
CFP prostheses								
0 to 1	44	0	44	0	44	0	100	100
1 to 2	44	0	44	0	44	0	100	100
2 to 3	44	0	44	0	44	0	100	100
3 to 4	44	0	44	2	43	0	100	100
4 to 5	42	0	42	2	41	2	95.3	95.3
5 to 6	38	0	38	0	38	1	97.4	92.8
6 to 7	37	0	37	0	37	0	100	92.8
FP prostheses								
0 to 1	141	0	141	2	140	0	100	100
1 to 2	139	0	139	3	137.5	0	100	100
2 to 3	136	0	136	2	135	2	98.5	98.5
3 to 4	132	0	132	3	130.5	3	98.7	97.2
4 to 5	126	0	126	4	124	2	98.4	95.6
5 to 6	120	0	120	4	118	0	100	95.6
6 to 7	116	0	116	2	115	0	100	95.6
FC prostheses								
0 to 1	33	0	33	0	33	0	100	100
1 to 2	33	0	33	0	33	0	100	100
2 to 3	33	0	33	6	30	0	100	100
3 to 4	27	0	27	6	24	0	100	100
4 to 5	21	0	21	0	21	0	100	100
5 to 6	21	0	21	0	21	0	100	100
6 to 7	21	0	21	0	21	0	100	100
ITS prostheses								
0 to 1	24	0	24	0	24	0	100	100
1 to 2	24	0	24	0	24	0	100	100
2 to 3	24	0	24	0	24	0	100	100
3 to 4	24	0	24	0	24	1	95.9	95.9
4 to 5	23	0	23	1	22.5	1	95.6	91.7
5 to 6	21	0	21	1	20.5	0	100	91.7
6 to 7	20	0	20	0	20	0	100	91.7
OD prostheses								
0 to 1	42	0	42	0	42	0	100	100
1 to 2	42	0	42	0	42	0	100	100
2 to 3	42	0	42	2	41	0	100	100
3 to 4	40	0	40	0	40	0	100	100
4 to 5	40	0	40	0	40	0	100	100
5 to 6	40	0	40	0	40	0	100	100
6 to 7	40	0	40	0	40	3	92.5	92.5

Table 10 Life Table Analysis for Survival of Implants Supporting ODs

Interval (y)	Implants at start of interval	Early failures	Loaded implants	Dropouts	Implants at risk	Failures during interval	Survival rate (%)	Cumulative survival rate (%)
3 or more implants supporting ODs								
0 to 1	92	0	92	0	92	0	100	100
1 to 2	92	0	92	0	92	0	100	100
2 to 3	92	0	92	2	91	0	100	100
3 to 4	90	0	90	0	90	0	100	100
4 to 5	90	0	90	0	90	0	100	100
5 to 6	90	0	90	0	90	0	100	100
6 to 7	90	0	90	0	90	3	96.7	96.7
2 implants supporting ODs								
0 to 1	34	0	34	0	34	0	100	100
1 to 2	34	0	34	2	33	0	100	100
2 to 3	32	0	32	0	32	0	100	100
3 to 4	32	0	32	2	31	2	93.6	93.6
4 to 5	28	0	28	4	26	0	93.6	93.6
5 to 6	24	0	24	2	23	0	93.6	93.6
6 to 7	22	0	22	0	22	0	93.6	93.6

for the application of short implants to the posterior mandible was suggested. In contrast to the aforementioned studies, Fartash and colleagues⁶⁰ documented a lower success rate for short implants than for long implants. In addition, Bert⁶¹ suggested the use of wide-diameter implants with tapping placement. Although the clinical results of short implants were favorable, further research is required to elucidate the most appropriate implant size and distribution, as well as the most favorable prosthetic restorations.

CONCLUSIONS

Clinical and radiographic preliminary results have demonstrated that restoration of complete and partially edentulous arches with ITI implants can be a reliable medium- to long-term treatment in this study population.

Medium- to long-term implant survival and success were not influenced by position site (maxilla or mandible). More research is required to explain the influence of implant placement sector (anterior or posterior) on implant survival and success.

Implants supporting fixed ST prostheses, CFP prostheses, FP prostheses, and ITS prostheses demonstrated similar 7-year survival rates. OD survival rates approximated those for FC prostheses, although ODs were more susceptible to prosthetic

complications than other types of prostheses. Nevertheless, more research is needed to confirm this trend, owing to the reduced number of implants supporting this type of prosthesis. Implant and prosthetic survival rates for ODs supported by 2 implants were comparable to those for ODs supported by 3 or more implants.

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