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Conflicts of interest

The authors declare no conflict of interest.

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A systematic review on survival and success rates of implants placed immediately into fresh extraction sockets after at least 1 year

Key words: extraction sockets, immediate implants, implant dentistry, success, survival, systematic review

Abstract

Background: Type I immediate implant placement has gained popularity because it may reduce treatment time, number of surgeries and post-extraction bone loss. However, this is potentially challenged by inadequate keratinized mucosa for flap adaptation and difficulties in achieving primary stability. Moreover, it has been proven that post-extraction bone loss is an inevitable biological process, which affects treatment outcomes.

Objectives: To estimate survival and success rates of implants and the implant-supported prostheses, the prevalence of biological, technical and aesthetic complications, and the magnitude of soft and hard tissue changes following implant placement immediately into fresh extraction sockets.

Material and methods: An electronic search in MEDLINE (PubMed) and the Cochrane Library from 1991 to July 2010 was performed to include prospective studies on immediate implants with a mean follow-up time of at least 1 year. The survival rates were computed using the STATA statistical software. Weighted means of soft and hard tissue changes were obtained by the inverse variance method.

Results: A total of 46 prospective studies, with a mean follow-up time of 2.08 years, were included. The annual failure rate of immediate implants was 0.82% (95% CI: 0.48–1.39%), translating into the 2-year survival rate of 98.4% (97.3–99%). Among the five factors analysed (reasons for extraction, antibiotic use, position of implant [anterior vs. posterior, maxilla vs. mandible), type of loading], only the regimen of antibiotic use affected the survival rate significantly. Lower failure rates were found in groups that were provided with a course of post-operative antibiotics. The success of implant therapy was difficult to assess due to scarce reporting on biological, technical and aesthetic complications. Soft tissue changes occurred mostly in the first 3 months after the provision of restoration, and then stabilized towards end of the first year. Marginal bone loss predominantly took place in the first year after implant placement, with a magnitude generally less than 1 mm. Controversy on hard tissue preservation with platform-switching technique remained unsolved.

Conclusions: Despite the high survival rate observed, more long-term studies are necessary to determine the success of implant treatment provided immediately after tooth extraction. Special attention has to be given to aesthetic outcomes.

Forty years ago, the first dental implant to replace a missing tooth in human oral cavity was reported (Brånemark et al. 1969). It was a sensational breakthrough in dentistry as it marked a new era to restore chewing function and aesthetics. Ever since, implant dentistry developed emphasizing aspects like dental materials, surface chemistry (Jansen et al. 1991; Klokkevold et al. 1997; Lazzara et al. 1999; Salvi et al. 2004, surface characteristics (Carlsson et al. 1988); Buser et al. 1991; Abrahamsson et al. 2004), as well as soft and hard tissue biology. The technique of placing titanium oral implants in healed edentulous sites and subsequently restoring the implant with a prosthesis has been recognized to be a highly predictive treatment for fully and partially edentulous patients. In general, the 5-year survival rate of implants is approximately 95%, and the 10-year survival rate is greater than 89% (Pjetursson et al. 2004). Nevertheless, over the years, researchers tried to minimize the treatment time needed and hence, timing of implant placement has recently drawn a large share of attention.

As the debate in timing of implant placement increased, the following new classification based on morphologic, dimensional and histologic changes that follow tooth extraction was proposed at the Third ITI Consensus Conference (Hämmerle et al. 2004):

- Type 1: Immediate placement: an implant is placed immediately in an extraction socket as part of the same procedure with no healing of bone or soft tissues.
- Type 2: Early placement (typically 4– 8 weeks of healing) with some soft tissue healing: the post-extraction site has healed soft tissue coverage of the alveolus but without significant bone healing.
- Type 3: Early placement with partial bone healing (typically 12–16 weeks of healing): The post-extraction site has both healed soft tissues and a significant degree of bone healing.
- Type 4: Late placement (more than 6 months after extraction): implant placement in a fully healed edentulous site.

Previously, practitioners allowed a socket healing time of 12 months or longer before placing dental implants to restore an edentulous space (Adell et al. 1981). Such a lag time brought the patient compromised comfort, function, and aesthetics. In 1978, the first report of a situation, in which the extraction followed by the placement of an implant into the fresh socket at the same appointment, was described as the "Türbingen immediate implant" (Schulte et al. 1978). This method reduced the number of dental appointments, the time of treatment and the number of surgeries required.

While immediate implant placement may yield attractive advantages, it has inherent disadvantages. The potential lack of keratinized mucosa for flap adaptation makes primary closure more difficult to be achieved in Type 1 than other types of implant placement. Moreover, the incongruity of size and shape between implants and extraction sockets presents challenges to primary implant stability. While initial implant stability is obtained by intimate contact with the pristine bone in healed sites, residual bony defects always exist around implants in Type 1 immediate implantation. Consequently, primary stability is only achieved by anchoring the implant in the apical bony region (3

-4 mm), where cancellous bone predominates. Moreover, although both animal (Araujo et al. 2005) and human studies (Covani et al. 2004a) show that spontaneous bone fill occurs in the peri-implant marginal defects after 3-4 months when the defect size is 2 mm or less, immediate implant placement cannot prevent intra- and extra-alveolar modelling and remodelling leading to the inevitable vertical and horizontal reduction in both buccal and lingual alveolar bony walls, conspicuously in the buccal aspect. Such biological changes imply higher risk of marginal mucosal recession after immediate implant placement, and hence, non-aesthetic restorations in areas of aesthetic priority may result, especially when the facial socket wall and tissue biotype are thin (De Rouck et al. 2008b).

In the posterior region, immediate implants may also be exposed to the dilemma between the difficulty in achieving primary stability if placed in the centre of the socket or a substantial defect if positioned leaning towards either wall of the socket. Consequently, more questions are invited such as the minimum dimension of defects in need of grafting (Botticelli et al. 2003), subsequently the choice of grafting material as well as the risk in graft exposure and management. All these aspects related to immediate implants could possibly lower the survival rate of the implant.

In a series of systematic reviews (Pjetursson et al. 2004, 2007; Jung et al. 2008), survival and complication rates of fixed dental prosthesis, formerly "fixed partial dentures" (FPDs) and single crowns supported by oral implants were estimated. The 5-year survival rate of implants was >95% and that of FPDs and SCs were approximately 95%. Technical and biological complications were reasonably prevalent. In the review concerning peri-implant diseases (Zitzmann & Berglundh 2008), it was found that after 5 -10 years in function, peri-implant mucositis occurred in approximately 80% of the subjects and in 50% of the implants. Peri-implantitis was found in 28-56% of the subjects and 12-43% of the implants.

In a recent Cochrane systematic review, success, complications, aesthetics and patient satisfaction among different timing of implant placement (immediate, immediate-delayed and delayed) after tooth extraction were evaluated (Esposito et al. 2010). Two studies of parallel group design, comparing immediate and delayed implant placement were included in this review. The meta-analysis of the two trials did not show any statistically significant difference between the two groups regarding prosthesis and implant failures. Concerning immediate vs. immediate-delayed implant placement only one trial was included in the review. There were eight patients in each group. Two years after implant placement, no implant failure and complications occurred, and no statistically significant difference was found with respect to the level of periimplant marginal mucosa and marginal bone level changes. Based on the few under-powered trials, it was concluded that there was insufficient evidence to determine possible advantages or disadvantages of immediate, immediate-delayed or delayed implants (Esposito et al. 2010).

Therefore, the main objectives of this systematic review are to quantitatively estimate the survival and success rates of immediate implants and the implant-supported prosthesis, the prevalence of biological, technical and aesthetic complications, and the magnitude of soft and hard tissue changes following implant placement in fresh extraction sockets (Type 1).

Material and methods

Search strategy

An electronic search in MEDLINE (PubMed) and the Cochrane library from January 1991 to July 2010 was performed using the following search terms:

{Intervention}

[immediate implant*] OR [immediate implant placement*] OR [(immediate implant*] AND (extraction socket*)] OR [immediate implant installation*] OR [early implant placement*] OR [early implant installation]

and in combination with the outcome terms: AND

{Outcome}

[survival] **OR** [complication*] **OR** [failure*] Moreover, manual searches of the biliographies of all full text articles and the following journals from January 2000 to December 2010 were also conducted:

- Clinical Oral Implants Research
- International Journal of Oral & Maxillofacial Implants
- Journal of Clinical Periodontology
- Journal of Periodontology
- Journal of Prosthetic Dentistry
- Clinical Implant Dentistry & Related Research
- International Journal of Periodontics & Restorative Dentistry
- International Journal of Prosthodontics.

Inclusion criteria

The studies to be selected had to yield the following inclusion criteria:

- English publications in the dental literature
- Human adults (≥18 years old) in good general health
- RCTs, prospective cohort studies and case series with a minimum of 10 subjects in the immediate implant placement group
- Studies with a mean follow-up time > 12 month following implant placement
- Studies reporting survival rates of the immediate implants
- Studies without multiple interventions (e. g. sinus augmentation via the transalveolar approach)
- Studies which clearly state timing of restorations or loading protocol.

Selection of studies

After the electronic search, two independent reviewers (K. Y. L. and L. P.) screened all titles and determined the number of abstracts to be evaluated. Following this, the two independent reviewers screened all selected abstracts for possible inclusion in the review and determined the selection of full-text articles. The full texts of all studies of possible relevance were then obtained for independent assessment by the reviewers. Any disagreement was resolved by discussion. The κ -values were 0.85 and 0.76 at the title and abstract levels, respectively.

Fig. 1 describes the process of identifying the 46 studies selected from an initial yield of 5887 titles. In the included studies, two had publications repeated on the same patient cohorts. In this situation, only the one with a longer observation period was chosen. Reasons for exclusion of articles not considered were noted as well.

Excluded studies

Of the 164 full-text articles examined, 118 were excluded from the final analysis (reasons, see reference list).

The main reasons for exclusion were: (Fig. 1):

- Not reporting on immediate implants,
- Unknown survival rate of immediate implants in the study (no report/no separate report of the survival rate of immediate implants from other types of implant insertion),
- Unknown number of immediate implants,
- Mean follow-up time less than 1 year or unknown mean follow-up time,

- The sample size (number of subjects) less than 10 in the immediate implant group,
- Unknown number of patients treated with immediate implants,
- Multiple interventions (e.g. Sinus lift augmentation via the transalveolar approach) were carried out simultaneously with immediate implants and
- Unknown timing for restorations or unknown loading protocol.

Quality assessment

Two reviewers (K. Y. L. and L. P.) independently assessed the quality of randomized controlled trials and prospective cohort studies.

Randomized controlled trials

The risk of bias of RCTs was assessed according to the recommended approach suggested by the Cochrane Collaboration. In this twopart tool, six specific domains were addressed, namely sequence generation, allocation concealment, blinding, incomplete outcome data, selective outcome reporting and other issues. A judgement of "Yes" indicated low risk of bias, "No" indicated high risk of bias, and "Unclear" indicated unclear or unknown risk of bias.

An RCT was assigned "Low risk of bias" if all key domains were of low risk of bias, "Unclear risk of bias" if there was unclear risk of bias of ≥ 1 key domains, and "High risk of bias" if ≥ 1 domains belonged to high risk of bias.

Prospective cohort studies

The quality of prospective cohort studies will be assessed using the Newcastle–Ottawa Scale (Wells et al. 2009). A maximum of nine



Fig. 1. Search strategy.

stars could be given to each study. Any study that scored less than five stars was excluded.

Data extraction

Data were extracted independently by the two reviewers (K. Y. L. & L. P.) using a data extraction form. Disagreement regarding data extraction was resolved by consensus after discussion.

Of the 46 studies included, information on the survival of the implants was retrieved. Survival was defined as implants remaining *in situ* at the follow-up examinations, irrespective of their conditions. Failure was defined as implants that were lost after implant placement immediately into the extraction socket.

For studies with a mean follow-up time of longer than 3 years, information regarding biological, technical and aesthetic complications was also extracted.

Biological complications included periimplant mucositis and peri-implantitis. Peri-implant mucositis was defined as the presence of inflammation in the mucosa at the implant with no signs of supporting bone loss. Peri-implantitis was defined as the presence of inflammation in the mucosa and loss of supporting bone at the implant (Zitzmann & Berglundh 2008).

Technical complications denoted mechanical damage to implants, to implant components and/or the suprastructures. They included fractures of the implants, loss of retention, screw/abutment loosening, loss of access hole restorations, fracture of abutments/screws and fracture of veneering materials/framework of prosthesis.

Aesthetic outcomes were assessed by the Pink Esthetic Score (PES) introduced by Furhauser et al. (2005), and/or the papilla index described by Jemt (1997).

The PES was based on seven parameters: mesial papilla, distal papilla, soft tissue level, soft tissue contour, alveolar process deficiency, soft tissue colour, and texture. Each parameter was assessed with a 2-1-0 score, with 2 being the best and 0 being the worst result. A maximum score of 14 can be achieved. The aesthetic outcome was optimal if the PES was ≥ 10 .

The papilla index (Jemt 1997) described the fullness of papillary fill:

- Index 0 = no papilla present
- Index 1 = less than one half the papilla height is present and a convex nature of the adjacent tissue nature is noted.
- Index 2 = greater than half the height of the papilla is present but not to the full

extent of the contact point. Papilla is not in complete harmony.

- Index 3 = the papilla fills the entire proximal space and is in good harmony.
- Index 4 = the papilla is hyperplastic.

Data regarding marginal soft tissue changes in the vertical dimension and radiographic bony changes after immediate implant placement were also extracted.

Statistical analysis

Failure rates were calculated by dividing the number of events (failure or complication) in the numerator by the total exposure time (implant-time) in the denominator.

The numerator was usually extracted directly from the publications. The total exposure time was calculated by taking the sum of

- 1. The exposure time of implants that could be followed for the whole observation time.
- 2. The exposure time up to a failure of the implant that was lost during the observation time.
- 3. The exposure time up to the end of observation time for implants that did not complete the observation period due to reasons such as death, change of address, refusal to participate in the follow-up, chronic illness, missed appointments and work commitments.

For each study, event rates for implants were calculated by dividing the total number of events by the total implant exposure time in years. For further analysis, the total number of events was considered to be *Poisson* distributed for a given sum of implant exposure years and *Poisson* regression with a logarithmic link-function and total exposure time per study as an offset variable were used (Kirkwood & Sterne 2003a).

Robust standard errors were calculated to obtain 95% confidence intervals (CIs) of the summary estimates of the event rates. The Spearman goodness-of-fit statistics and associated P-values were calculated to assess heterogeneity of the study specific event rates. If the goodness-of-fit P-value was below 0.05, indicating heterogeneity, random-effects Poisson regression (with Gamma-distributed random-effects) was used to obtain a summary estimate of the event rates. One-year survival proportions were calculated via the relationship between event rates and survival function S, $S(T) = \exp(-T \times \text{event rate})$, by assuming constant event rates (Kirkwood & Sterne 2003b). The 95% CI for the survival proportions were calculated by using the 95% confidence limits of the event rates.

Multivariable *Poisson* regression was used to investigate whether reasons for extractions, use of antibiotics, locations of implants (anterior vs. posterior, maxillary vs. mandibular), and timing of restorations would affect the survival rate of immediate implants.

All analysis were performed using STATA/ SE[®], version 11 (Stata Corp., College Station, TX, USA).

To calculate vertical marginal soft tissue changes and radiographic bony changes, data obtained from studies at a specific time interval after restorations on immediate implants were pooled to derive the weighted mean and variance by the inverse variance method. The weighted mean was:

$$\bar{x} = \frac{\sum_{i=1}^{n} (x_i / \sigma_i^2)}{\sum_{i=1}^{n} (1 / \sigma_i^2)},$$

and the variance of the weighted mean was:

$$\sigma_{\bar{x}}^2 = \frac{1}{\sum_{i=1}^n (1/\sigma_i^2)},$$

where x_{i} , σ_i^2 were the known mean and variance obtained from each study.

Moreover, to test for the homogeneity of the results of the studies, which contributed to a particular weighted mean, the chi-squared (χ^2 , or Chi²) test among these studies was carried out:

$$Q = \sum w_i(x_i - \bar{x})$$

where Q was the chi-squared statistic; w_i was the reciprocal of the variance of the effect of the *i*th study, i.e. $1/\sigma_i^2$; \bar{x} was the weighted mean; and x_i was the mean of the *i*th study. To quantify the inconsistency across the studies, the following statistic was used:

$$I^2 = \left(\frac{Q - df}{Q}\right) \times 100\%,$$

where df was its degrees of freedom (i.e. df = i-1, where *i* was the number of studies).

A rough guide to interpretation was as follows:

- $I^2 = 0\%$ to 40%: might not be important;
- I² = 30% to 60%: may represent moderate heterogeneity;
- I² = 50% to 90%: may represent substantial heterogeneity;
- I² = 75% to 100%: considerable heterogeneity.

Results

A total of 46 prospective studies on implants inserted immediately into extraction sockets were included in this systematic review.

With the exception of three studies (Becker et al. 1994; Lang et al. 1994; Becker et al. 1998), all the other 43 studies were published after the year 2000. Most studies had the mean follow-up time less than 3 years. Only nine studies reported implant survival rates with the mean observation period of 3 years or more.

The studies were mainly conducted in an institutional environment. Five studies were multicenter studies. There were 16 comparative studies. The test and control groups were recruited to compare (i) implants placed in extraction sockets vs. implants inserted at healed sites (Kan et al. 2007a; Ribeiro et al. 2008; Siciliano et al. 2009; Gokcen-Rohlig et al. 2010); (ii) immediate implants at sites with chronic periapical lesion vs. implants at healed sites (Lindeboom et al. 2006a); (iii) immediate implant placement at sites with chronic periapical lesion vs. at sites without periapical lesion (Crespi et al. 2010); (iv) implants in acutely infected sockets vs. implants in sockets without pathology (Siegenthaler et al. 2007); (v) immediate implant insertion and simultaneous connective tissue graft vs. coronally advanced flap (Cornelini et al. 2008); (vi) treatment of immediate implant and connective tissue graft vs. immediate implant only (Bianchi & Sanfilippo 2004); (vii) immediate implantation with GBR vs. without GBR (Bragger et al. 1996); (viii) submerged vs. non-submerged healing following implant placement in extraction sockets (Cordaro et al. 2009); (ix) immediate vs. delayed provisional restoration after immediate implant placement (Crespi et al. 2008; De Rouck et al. 2009; Prosper et al. 2010); (x) immediate implants restored with a platform-switching vs. a platform matching protocol (Crespi et al. 2009a; Canullo et al. 2009a,b).

Ten studies were randomized clinical trials (Bianchi & Sanfilippo 2004; Lindeboom et al. 2006a; Crespi et al. 2008; Cornelini et al. 2008; Cordaro et al. 2009; De Rouck et al. 2009; Canullo et al. 2009a,b; ; Crespi et al. 2009a; Siciliano et al. 2009; Prosper et al. 2010). Each study was assessed according to the recommended approach suggested by the Cochrane Collaboration. Four studies were judged to be at high and the remaining six studies at unclear risk of bias. (Fig. 2 and Table 1)

Five studies were prospective cohort studies, and were assessed using the Newcastle– Ottawa Scale (Wells et al. 2009). All of them scored eight of nine stars.

A total of 2130 patients, aged between 18 and 94 were included in the 46 studies. Totally, 3082 implants were placed in which 2934 were in fresh extraction sockets and 148 were in healed sites. Twenty-five implants in two studies were not restored (Fugazzotto 2002a); Vidal et al. 2010), and one implant in one study was lost to followup (Calvo-Guirado et al. 2009), leaving 2908 implants for further analysis. (Table 2)

Reasons for extraction

In 12 studies, implants were used to replace teeth extracted due to non-periodontal reasons, e.g. root fractures, caries, endodontic failure and root resorption (Table 2). Removal of teeth because of both periodontal and nonperiodontal reasons was reported in 21 studies. In 13 studies, reasons for extraction were unclear.

Antibiotic prophylaxis

Antibiotics were prescribed in 33 studies (Table 2). Four studies involved pre-operative single dose of antibiotic prophylaxis, while post-operative antibiotic use of 5– 7 days was reported in 15 studies. Fourteen studies provided both pre-operative single dose and post-surgical (5–7 days) antibiotic prescription. Data regarding antibiotic prescription were not forthcoming in the remaining studies.

Position of implants – anterior vs. posterior

In five studies, implants were inserted in the anterior region only, namely, central incisors, lateral incisors and canines (Table 2). Implants in another five studies were solely placed in posterior areas, i.e. premolars and molars. Thirty-five studies involved implantation in both anterior and posterior regions. One study did not state the implant locations.

Position of implants - maxilla vs. mandible

Eighteen studies reported on implants installed in the maxilla only, while only three studies had all implants placed in the mandible (Table 2). In 23 studies, both maxillary and mandibular arches were involved, and two studies did not explicitly specify the arch that implants were inserted.

Types of implants

Various implant systems were employed in the 46 studies (Table 2). The majority of the implants were rough surface implants. Only three studies used implants of machined surfaces (Goldstein et al. 2002; Becker et al. 1994, 1998).

Grafting materials

No grafting materials were utilized at all in six studies (Table 2). Eleven studies involved autogenous bone grafts. Bone substitutes were used in 16 studies, of which demineralised bovine bone matrix (DBBM) was most frequently applied. Other reported bone substitutes were demineralised freeze-dried bone allograft (DFDBA), enamel matrix derivatives (EMD), Biogran[®] bone graft, and HRT synthetic bone allograft. The main purposes of using grafting materials were to fill the marginal gaps between implants and socket walls and to cover bony dehiscences and/or fenestrations.



Fig. 2. "Risk of bias graph" of included RCTs.

Table 1.	Summary	of risk	of bias	of	included	RCTs
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	Adequate sequence generation?	Allocation concealment?	Blinding?	Incomplete outcome data addressed?	Free of selective reporting?	Free of other bias?
Prosper et al. 2010	Unclear	Unclear	High	Low	Low	Low
Cordaro et al. 2009	Low	Low	High	Low	Low	low
De Rouck et al. 2009	Low	Unclear	Low	Low	Low	Low
Crespi et al. 2009a	Unclear	Unclear	High	Low	Low	Low
Canullo et al. 2009a,b	Low	Low	High	Low	Low	Low
Siciliano et al. 2009	High	High	High	Low	Low	Low
Crespi et al. 2008	Unclear	Unclear	Low	Low	Low	Low
Cornelini et al. 2008	Unclear	Unclear	Unclear	Low	Low	Low
Lindeboom et al. 2006a	Low	Unclear	Unclear	Low	Low	Low
Bianchi & Sanfilippo 2004	Unclear	Unclear	Low	Low	Low	Low

In 12 studies, the grafting materials were covered by barrier membranes. Resorbable membranes were more commonly used than non-resorbable membranes. In five studies, bony defects around implants were solely covered by barrier membranes.

Besides bone substitutes and barrier membranes, connective tissue grafting was also performed to cover immediate implants in six studies. Three studies reported a combined use of subepithelial connective tissue grafts (SCTG) with other grafting materials, while the other three employed SCTG as the only grafting material. The most common donor site was the palatal vault. This technique was mainly applied on subjects who were of the thin gingival biotype.

Loading

According to the Fourth ITI Consensus Report (Weber et al. 2009), immediate loading of dental implants was defined as loading being earlier than 1 week subsequent to implant placement; early loading was defined as loading being between 1 week and 2 months subsequent to implant placement; and conventional loading was loading being greater than 2 months subsequent to implant placement (Table 2).

Using definitions by this consensus report (Weber et al. 2009), implants in 19 studies were immediately loaded after implant insertion. Most of them used acrylic resin as the material for provisional restorations. Among the 19 studies, the provisionals were screwretained in six studies and cement-retained in nine studies. One study used both methods for retention. Implants in 23 studies were conventionally restored. The time of loading varied from 8 weeks to 1 year, but most of them were loaded within 3–6 months. The remaining four studies had both immediate and conventionally loading groups.

Definitive restorations

In most studies, implants placed immediately into extraction sockets were used to replace single missing teeth and hence, the most prevalent prostheses constructed were single crowns (Table 2). In seven studies, some implants also served as abutments for implant–implant or implant–tooth supported fixed dental prostheses. In more than half of the studies, permanent restorations were retained by cement, while screwretained restorations were used in only four studies.

Implant-supported overdentures were also reported in three studies (Huys 2001; Gokcen-Rohlig et al. 2010; Vidal et al. 2010). All the implants were conventionally loaded. Gokcen-Rohlig et al. (2010) used *locator* abutments, while Huys (2001) delivered ballretained overdentures. In the third study, the type of abutments was not specified (Vidal et al. 2010).

Survival of implants

Survival was defined as implants remaining *in situ* at the follow-up examinations, irrespective of their conditions. Failure was defined as implants that were lost after immediate implant placement.

The 46 included studies provided data on 2908 implants with the mean follow-up time of 2.08 years following implant placement into the extraction sockets. Fifty-eight implants were lost during the observation period. The estimated annual failure rate of the implants was 0.82% (95% CI: 0.48–1.39%), yielding the 2-year survival rate of 98.4% (97.3–99%) (Table 3).

When the nine studies with the mean follow-up time ≥ 3 years were analysed separately, the estimated annual failure rate was 0.62% (95% CI: 0.31–1.23%), translating into a 4-year implant survival rate of 97.5% (95.2– 98.8%) (Table 4).

Five factors were investigated for their impact on the survival of immediate implant: use of antibiotics, reasons for extractions, locations of implants (anterior vs. posterior, maxillary vs. mandibular), and timing of restorations:

Antibiotics

Four studies, with a total of 244 implants, involved a pre-operative single dose of antibiotic prophylaxis (Tables 5 and 6). Post-operative antibiotic use of 5-7 days was reported in 15 studies, involving 935 implants. Fourteen studies prescribed both pre-operative single dose and 5-7 days of post-operative antibiotics, and a total of 665 implants were examined. The relative failure rates of the three different groups were analysed with multivariable fixed-effect Poisson regression using pre-surgical antibiotic prophylaxis as the reference. The estimated annual failure rate for the pre-operative antibiotic use group was 1.87%. Both the post-operative antibiotic use group and pre- and post-operative antibiotic use group showed lower annual failure rates than did the pre-operative antibiotic use group, with the annual failure rates of 0.51% and 0.75%, respectively. The differences reached statistical significance (P = 0.002;0.02).

Reasons for extraction

In 12 studies, 424 implants were used to replace teeth extracted due to non-periodontal reasons (Table 7). Removal of teeth because of both periodontal and non-periodontal reasons was reported in 21 studies, involving 1094 implants. The estimated annual failure rate of the former group was 0.81%, and that of the latter group was 0.92%. The difference in the failure rates was tested by using a random-effect *Poisson* regression analysis, and this difference did not reach statistical significance (P = 0.84).

Position of implants - anterior vs. posterior

In the 46 studies included, a total of 486 implants were placed in the anterior region and 967 implants were inserted in the posterior region (Table 8). The locations of the remainder of the 1455 implants were unclear. The estimated annual failure rate of implants placed in the posterior area was slightly higher than that of those placed in the ante-

ndies
patients range age
n 51 23-94 59 c
n 12 22–54 Unknown
n 10 49–68 54
n 30 34-71 51.2
503 CL 3C 11 503
C.OC 21-02 1/ 11
30 18-70 Unknown hter
n 50 29-51 39.64
n 49 ≥ 18 IRG: 55,
ח 30 (15 ≥ 18 48.1 immediate = 18 48.1 immediate
n 45 25-67 48.73
n 20 28-71 52.3
n 30 31-75 55.8
22 32-76 50
n 26 19-76 60
n 16 23-62 42
n 46 18-71 47.2
n 30 24–76 54
n, 82 21–85 40.2 tter
a4 21-62 43

Restorations- definitives	Single crowns (cemented)	CMC (cemented)	Single crowns (unknown type of retention)	10 as abutments of FPDs (cemented), 11 single crowns	Single crowns (cemented/ screw-retained)	Single crowns (unknown twne of retention)	Single crowns (unknown tuno of retention)	type of retention) CMCs (cemented)	Single crowns (cemented)	9 single crowns, 15 partial, 11 complete (comanted)	CMCs (cemented)	Single crowns (cemented)	Single CMCs 6 months after implant nlacement	Single crowns (unknown type of retention)	Single crowns (unknown type of	Splinted FPDs (unknown type of retention)	Single crowns (unknown type of	CMCs/all ceramic crowns (cemented)
Restorations on II-Timing	All conventional	Immediate: 20, conventional: 20	All immediate	All conventional	All immediate	Immediate: 7, conventional: 22	All conventional	All immediate	All conventional	All immediate	All immediate	All conventional	All immediate	All conventional	All immediate	All immediate	All immediate	All immediate
Grafting materials	DBBM/ demineralized allograft + cortical chips + Ti-reinforced ePTFE membrane or bioabsorbable	Unknown	Unknown	No grafting materials	Autogenous bone graft + Bio-Oss	Bio-Oss + Bio-Gide	Connective tissue graft	Autogenous bone/ xenograft + BioGide, + Subepithelial connective	Bio-Oss + Bio-Gide + connective tiscua oraft	Autogenous bone chips	No grafting materials	Autogeneous corticocancellous	Autogenous bone grafts	EMD/bioabsorble collagen membrane	BioGide	Autogenous bone graft + resorbable membrane (Gore Resolut	Biogran bone graft 3i + Bio.Gide	Unknown
Implant	Straumann	Outlink	Conect, Conic	Straumann	TiUnite Nobel Biocare; Nobel Perfect Nobel Biocare	Straumann	Padova	Nobel Biocare: Replace select/ NobelPerfect	Nobel Biocare; Replace select	Outlink	Premium	Frialit 2 synchro	Frialit II	Straumann	Straumann TE	Mk IV TiUnite (Branemark)	NT Ossseotite 3I, Frialit-2	Astra Tech ST
Position - Mx vs. Md	All Md: 341	All Mx: 40	All Mx: 46	Mx: 16, Md: 5	All Mx: 23	Mx: 23, Md: 6	Mx: 7, Md: 3	All Mx: 23	All Mx: 14	Mixed	Mixed	Mixed	All Mx: 33	Mx: 15, Md: 17	Mx: 19, Md: 3	Mx: 39, Md: 11	All Mx: 28	All Mx: 16
Position - Ant vs. Post	All Post	Ant: 24, Post: 16	Mixed	Ant: 5, post: 16	Mixed	Ant: 14, Post: 15	Ant: 5, Post: 5	All Ant	All Ant	Mixed	Mixed	Mixed	Ant: 26, Post: 7	Ant: 15, Post: 17	Ant: 9, Post: 13	Ant 19, Post: 31	Mixed	Ant: 15, Post 1
Reasons for extraction	Unknown	Perio + non-nerio	Non-perio	Non-perio	Unknown	Non-perio	Perio + non-perio	Perio + non-perio	Non-perio	Unknown	Unknown	Unknown	Non-perio	Perio + non-perio	Perio + non-perio	Unknown	Unknown	Perio + non-nerio
Antibiotic use	Post	Pre + post	Post	Unknown	Post	Pre + post	Post	Post	Pre + post	Pre + post	Pre + post	Pre	Pre + post	Post	Post	Pre + post	Unknown	Pre + post
No of implants	341	40	82 (46 immediate implant)	21	38 (23 immediate implants)	29	10	33	14	160 (150 immediate implant)	18	50 (25 immediate imploate)	33	32	52	20	43 (28 immediate implants)	111. province, 28 (16 immediate
Mean age	Unknown	47.21	45.4	49.1	45.1	Test: 45; Control: 55	Unknown	39.5	28	57	Unknown	39.7	Unknown	45	39	55	Unknown	48.2
Age range	26-81	24-68	23-71	21-81	18-70	23-82	42-55	25-63	18-49	42-72	22–60	19–69	24-58	21–60	√ 18	35-66	20-60	27-72
No of patients	320	40	64	18	29 (19 immediate implants)	29	10	23	12	27	18	50 (25 immediate immediate	33	32	22	6	38	25 (16 immediate
Setting	Private	Institution	Institution & private	Private	Institution	Institution	Institution	Institution	Institution	Institution	Institution	Institution	Institution	Private	Unknown	Private	Institution	Private
Study design	Prosp.	Prosp., randomized	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.
Year of publication	2008	2008	2008	2008	2007	2007	2007	2007	2007	2007	2006	2006	2006	2005	2005	2005	2005	2004
Study	Fugazzotto	Crespi et al.	Ribeiro et al.	Botticelli	Kan et al.	Siegenthaler et al.	Covani et al.	Kan et al.	Juodzbalys et al.	Crespi et al.	Barone et al.	Lindeboom et al.	Ferrara et al.	Cangini and Cornelini	Cornelini et al.	Vanden Bogaerde et al.	Tsirlis	Norton

	Grafting Restorations Restorations- materials on II-Timing definitives	Autogenous All conventional CMCs (strew- bone graft + retained) biobscrabale membranes	SCTG All conventional CMCs (cemented)	Unknown All immediate CMCs (cemented)	Autogenous All conventional CMCs, FPDs bone chips + (cemented) gratting materials	DFDBA used to All conventional Single crowns fill the defect/ (unknown type of barrier retention) membranes	HRT synthetic All conventional 426: ball retained bone allograft All conventional 426: ball retained FPDs (cemented), 44 implant-tooth FPDs (cemented), 4 imgle crowns (cemented)	No grafting All conventional Unknown materials (unknown type of retention)	E-PTE barrier all conventional 13 CMCs membrane (11: cemented, 2: screw-retained), 3: unit FPDs, 4: abutmens of 4: abutmens of 4: abutment PPDs, 4: autoorted FPDs, 4: autoorted FPDs,	Oval 6 All conventional 11 single crowns, e-PTFE tissue 10 abutments for augmentation FPDs (cemented) material +
	Implant	Premium (Sweden & Martina)	Straumann	Replace, Nobel Biocare	Straumann/ Astra Tech	Branemark, 3I.	Straumann: hollow screws, solid screws.	Branemark	ITI Bonefit Dental implant System	Nobelpharma AB, Gothenburg, Sweden
	Position - Mx vs. Md	Mx: 95, Md: 68	Mixed	All Mx: 35	All Mx: 46	All Mx: 47	Unknown	Mixed	Mixed	Mixed
	Position - Ant vs. Post	Ant: 82, Post: 81	Mixed	All Ant: 35	All Post: 46	Ant: 16, Post: 31	Unknown	Mixed	Mixed	Mixed
	Reasons for extraction	Perio + non-perio	Perio + non-perio	Non-perio	Unknown	Unknown	Unknown	Perio + non-perio	Perio + non-perio	Perio + non-perio
	Antibiotic use	Post	Unknown	Post	Unknown J)	Unknown	Unknown	Pre	Post	Pre + post
	No of implants	163	115	35	63 (17 unrestored	47	556	134	28	49
	Mean age	Unknown	45.4	36.5	Unknown	45.79	Unknown	Unknown	Unknown	Unknown
	Age range	20-68	19–73	18-65	31–66	19–72	19–71	29-81	20-81	23-83
	No of patients	95	116	35	57	38	147	81	21	49
	Setting	Institution	Institution	Institution	Private	Institution	Unknown	Institution	Institution	Institution, multicenter
	Study design	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.	Prosp.
inued)	Year of publication	2004	2004	2003	2002	2002	2001	1998	1994	1994
Table 2. (conti	Study	Covani et al.	Bianchi et al.	Kan et al.	Fugazzotto et al.	Goldstein et al.	Huys	Becker et al.	Lang et al.	Becker et al.

rior area (0.54% vs. 0.45%). However, the difference was not statistically significant (P = 0.82).

Position of implants - maxilla vs. mandible

In total, 933 implants were inserted in the maxilla, and 731 implants in the mandible (Table 9). The remainders were not specified about the implant position. The implants placed in the maxilla had a higher estimated annual failure rate (0.73%) than implants placed in the mandible (0.50%). However, in the *Poisson* regression analysis, this difference was not statistically significant (P = 0.58).

Loading

In the included studies, much more implants were conventionally (n = 2086) than immediately loaded (n = 822) (Table 10). The estimated annual failure rate of the conventional loading group was lower than that of the immediate loading group (0.75% vs. 0.89%). However, the difference, again, did not reach statistical significance in the *Poisson* regression analysis (P = 0.73).

Success

With regard to the success of an implantrelated treatment, survival of the implants and their reconstructions should not be the ultimate goal of analysis. Rather, a successful treatment should be free of any biological and technical complications, and aesthetic outcomes should be satisfactory.

In this systematic review, the nine studies with the mean follow-up time of 3 years or more were evaluated for treatment success.

Biological complications

According to Lang & Berglundh (2011), the key parameter for the diagnosis of periimplant mucositis was bleeding on gentle probing; and peri-implantitis was characterized by changes in the level of the crestal bone in conjunction with bleeding on probing with or without concomitant deepening of peri-implant pockets. Pus was a common finding in peri-implantitis sites.

To investigate the biological complications, the following parameters were considered:

- 1. Bleeding and suppuration on probing
- Changes in marginal bone levels on radiographs

Among the nine studies, seven assessed radiographic bony changes; however, only three of them clinically assessed the periimplant soft tissue response to periodontal

Table 3. Annual failure rates and survival of implants inserted in extraction sockets

	Year of publication	Total no. of implants	Mean follow-up time	No. of failure	Before loading	After loading	Loss to follow-up	Total implant exposure time (years)	Estimated failure rate (per 100 implant years)	Estimated survival (%) after 2 years
Vidal et al.	2010	54	1	0	0	0	0	54	0	100
Tortamano et al.	2010	12	1.5	0	na	0	0	18	0	100
Gocen-Rohlig et al.	2010	20	2.33	0	0	0	0	46.67	0	100
Crespi et al.	2010	30	2	0	0	0	0	60	0	100
Prosper et al.	2010	120	5	4	1	3	0	580.46	0.69	98.62
Mijiritsky et al.	2009	24	3.27	1	na	1	0	78.5	1.27	97.45
Crespi et al.	2009	64	2	0	na	0	0	128	0	100
Kan et al.	2009	20	2.15	0	na	0	0	43	0	100
Del Fabbro et al.	2009	61	1.79	1	1	0	0	107.67	0.93	98.14
Canullo et al.	2009	22	2.08	0	na	0	0	45.83	0	100
Kahnberg	2009	40	2	0	0	0	0	80	0	100
Cordaro et al.	2009	30	1.5	1	1	0	0	43.73	2.29	95.43
Calvo-Guirado et al.	2009	61	1	1	na	1	1	59.33	1.69	96.63
De Rouck et al.	2009	49	1.07	3	2	1	0	52.33	5.73	88.54
Siciliano et al.	2009	15	1	0	0	0	0	15	0	100
Botticelli et al.	2008	21	5	0	0	0	0	105	0	100
Lops et al.	2008	46	1.15	0	0	0	0	53.08	0	100
De Rouck et al.	2008	30	1	1	na	1	0	29.08	3.44	93.12
Cornelini et al.	2008	34	1	0	na	0	0	34	0	100
Fugazzotto	2008	341	2.72	2	1	1	0	929.06	0.22	99.57
Crespi et al.	2008	40	2	0	0	0	0	80	0	100
Ribeiro et al.	2008	46	2.17	3	na	3	0	99.91	3	93.99
Cafiero et al.	2008	82	1	0	0	0	0	82	0	100
Crespi et al.	2007	150	1.5	0	0	0	0	225	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Siegenthaler et al.	2007	29	1	0	0	0	0	29	0	100
Covani et al.	2007	10	1	0	0	0	0	10	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Barone et al.	2006	18	1	1	na	1	0	17.08	5.86	88.29
Lindeboom et al.	2006	25	1	2	2	0	0	24	8.33	83.33
Ferrara et al.	2006	33	2.34	2	na	2	0	77.25	2.59	94.82
Cangini and Cornelini	2005	32	1	0	0	0	0	32	0	100
Cornelini et al.	2005	22	1	0	na	0	0	22	0	100
Vanden Bogaerde et al.	2005	50	1.5	0	0	0	0	75	0	100
Tsirlis	2005	28	2	0	0	0	0	56	0	100
Norton	2004	16	1.61	0	na	0	0	25.75	0	100
Covani et al.	2004	163	4	5	2	3	0	636.5	0.79	98.43
Bianchi et al.	2004	115	5.38	0	0	0	0	618.5	0	100
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Goldstein et al.	2002	47	3.37	0	0	0	0	158.5	0	100
Fugazzotto et al.	2002	46	1.40	0	0	0	0	64.33	0	100
Huys	2001	556	7	19	19	0	0	3763.75	0.50	98.99
Becker et al.	1998	134	3.61	9	2	7	0	483.17	1.86	96.27
Lang et al.	1994	28	3.28	0	0	0	0	91.93	0	100
Becker et al.	1994	49	1.51	3	3	0	0	74.06	4.05	91.90
Summary estimate (95%	CI) [*]	2908	2.08	58	34	24	1	9431.17	0.82 (0.48–1.39)	98.4
										(97.3–99)
*Based on random-effect	s Poisson rear	ession, test for	heterogene	itv <i>P</i> < 0.0)1.					

probing (Bianchi & Sanfilippo 2004; Botticelli et al. 2008; Prosper et al. 2010).

Bleeding and suppuration on probing

Three studies reported on the effectiveness of patients' self-performed oral hygiene (Bianchi & Sanfilippo 2004; Botticelli et al. 2008; Prosper et al. 2010). The prevalence of plaque accumulation on implants varied among the studies. In both studies of Prosper et al. (2010) and Botticelli et al. (2008), the sites with plaque accumulation were less than 20%. However, 40% of implant sites harboured plaque in the study of Bianchi & Sanfilippo (2004). As a result, BOP was more prevalent at implants in the latter study than in the two former studies (31% vs. 6-17%).

With the criteria defined above (Lang & Berglundh (2011), 31% of implants demonstrated peri-implant mucositis in the study by Bianchi & Sanfilippo (2004), while in the two other studies (Botticelli et al. 2008; Prosper et al. 2010), peri-implant mucositis was less prevalent.

Change in marginal bone levels

Seven studies with a mean follow-up time ≥ 3 years evaluated marginal bony alterations (Becker et al. 1998; Huys 2001; Bianchi & Sanfilippo 2004; Covani et al. 2004b;

Botticelli et al. 2008; Mijiritsky et al. 2009; Prosper et al. 2010). With the exception that there was a 0.23 mm gain in the mean radiographic bone level in one study (Botticelli et al. 2008), immediate implants in most studies experienced marginal bone loss after being in service. However, in most of the cases, the loss was within the range that fulfilled one of the success criteria stated by Albrektsson et al. (1986), namely, that "after the first year of service, the annual vertical bone loss should not exceed 0.2 mm."

Covani et al. (2004b) reported that 4 years after implant placement immediately into

Table 4. Annual failure rates and survival of implants inserted in extraction sockets. (studies with mean follow-up time \geq 3 years)

Study	Year of publication	Total no. of implants	Mean follow-up time	No. of failure	Before Loading	After loading	Loss to follow-up	Total implant exposure time (years)	Estimated failure rate (per 100 implant yeaers)	Estimated survival (%) after 4 years
Prosper et al.	2010	120	5	4	3	1	0	580.46	0.69	97.24
Mijiritsky et al.	2009	24	3.27	1	na	1	0	78.5	1.27	94.90
Botticelli et al.	2008	21	5	0	0	0	0	105	0	100
Covani et al.	2004	163	4	5	2	3	0	636.5	0.79	96.86
Bianchi et al.	2004	115	5.38	0	0	0	0	618.5	0	100
Goldstein et al.	2002	47	3.37	0	0	0	0	158.5	0	100
Huys	2001	556	7	19	19	0	0	3763.75	0.50	97.98
Becker et al.	1998	134	3.61	9	6	3	0	483.17	1.86	92.55
Lang et al.	1994	28	3.28	0	0	0	0	91.93	0	100
Summary		1208	4.43	38	30	8	0	6516.31	0.62	97.5
estimate (95% CI) [*]									(0.31–1.23)	(95.2–98.8)
*Based on rando	m-effects Poiss	on regressio	n, test for heterog	eneity P <	0.01.					

Table 5. Comparison of annual failure rates and survival of implants inserted in extraction sockets – antibiotic uses (pre-surgical vs. post-surgical vs. pre- and post-surgical)

Study	Year of publica-tion	Total no. of implants	Mean follow-up time (years)	No. of failure at 1 year after implant placement	Before loading	After loading	Lost to follow-up	Total implant exposure time (years)	Estimated failure rate (per 100 implant years)	Estimated survival after 2 years (%)
Pre-surgical antibiotic use										
Del Fabbro et al.	2009	61	1.79	1	1	0	0	107.67	0.93	98.14
Mijiritsky et al.	2009	24	3.27	1	na	1	0	78.5	1.27	97.45
Lineboom et al.	2006	25	1	2	2	0	0	24	8.33	83.33
Becker et al.	1998	134	3.61	9	2	7	0	483.17	1.86	96.27
Summary estimate (95%	CI)*	244	2.42	3.25	5	8	0	693.33	1.87	96.3 (93 7 97 8)
Post-surgical antibiotic use									(1.05-5.25)	(55.7-57.6)
Calvo-Guirado et al.	2009	61	1	1	na	1	1	59.33	1.69	96.63
Siciliano et al.	2009	15	1	0	0	0	0	15	0	100
Kan et al.	2009	20	2.15	0	na	0	0	43	0	100
Cafiero et al.	2008	82	1	0	0	0	0	82	0	100
Fugazzotto	2008	341	2.72	2	1	1	0	929.06	0.22	99.57
Ribeiro et al.	2008	46	2.17	3	na	3	0	99.91	3	93.99
Cornelini et al.	2008	34	1	0	na	0	0	34	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Covani et al.	2007	10	1	0	0	0	0	10	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Cangini and Cornelini	2005	32	1	0	0	0	0	32	0	100
Cornelini et al.	2005	22	1	0	na	0	0	22	0	100
Covani et al.	2004	163	4	5	2	3	0	636.5	0.79	98.43
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Lang et al.	1994	28	3.28	0	0	0	0	91.93	0	100
Summary estimate (95%	CI)*	935	1.65	11	3	8	1	2149.44	0.51	99
Pre + post-surgical antibiot	ic use								(0.15-1.97)	(90.1–99.7)
Prosper et al.	2010	120	5	4	1	3	0	580.46	0.69	98.62
Crespi et al.	2010	30	2	0	0	0	0	60	0	100
Crespi et al.	2009	64	2	0	na	0	0	128	0	100
Canullo et al.	2009	22	2.08	0	na	0	0	45.83	0	100
Crespi et al.	2008	40	2	0	0	0	0	80	0	100
De Rouck et al.	2008	30	1	1	na	1	0	29.08	3.44	93.12
Siegenthaler et al.	2007	29	1	0	0	0	0	29	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Crespi et al.	2007	150	1.5	0	0	0	0	225	0	100
Barone et al.	2006	18	1	1	na	1	0	17.08	5.86	88.29
Ferrara et al.	2006	33	2.34	2	na	2	0	77.25	2.59	94.82
Vanden Bogaerde et al.	2005	50	1.5	0	0	0	0	75	0	100
Norton	2004	16	1.61	0	na	0	0	25.75	0	100
Becker et al.	1994	49	1.51	3	3	0	0	74.06	4.05	91.90
Summary estimate		665	1.90	11	4	7	0	1474.51	0.75	98.5
(95% CI)*									(0.19–1.75)	(96.6–99.6)

*Based on fixed-effects *Poisson* regression, test for heterogeneity P = 0.1

Table 6. Summary	of annual failure rates,	relative failure rates and	d survival estimates for im	plants inserted in extraction	sockets - antibiotic uses
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Type of antibiotics use	Total number of implants	Total implant exposure time (years)	Mean follow-up time	Estimated annual failure rate [†]	2 year survival summary estimate (95% CI) [†]	Relative failure rate [*]	<i>P</i> -value [*]
Pre-surgical	244	693.33	2.42	1.87 (1.09–3.23)	96.3 (93.7–97.8)	1 (Ref.)	
Post-surgical	935	2149.44	1.65	0.51 (0.13–1.97)	99 (96.1–99.7)	0.27 (0.12–0.61)	0.002
Pre- and post- surgical	665	1474.51	1.9	0.75 (0.19–1.75)	98.5 (96.6–99.6)	0.40 (0.18–0.89)	0.024
*Based on multivariab	le fixed-effect Poi	isson regression.					

on fixed-effects *Poisson* regression

Table 7. Comparison of annual failure rates and survival of implants inserted extraction sockets - reasons for extraction

Study	Year of publication	Total no. of implants	Mean follow-up time (year)	No. of failure at 1 year after implant placement	Before loading	After loading	Lost to follow-up	Total implant exposure time (years)	Estimated failure rate (per 100 implant years)	Estimated survival after 2 year (%)
Non-periodontal reaso	ns									
Tortamano et al.	2010	12	1.5	0	na	0	0	18	0	100
Crespi et al.	2010	30	2	0	0	0	0	60	0	100
Calvo-Guirado et al.	2009	61	1	1	na	1	1	59.33	1.69	96.63
Siciliano et al.	2009	15	1	0	0	0	0	15	0	100
Lops et al.	2008	46	1.15	0	0	0	0	53.08	0	100
Cafiero et al.	2008	82	1	0	0	0	0	82	0	100
Ribeiro et al.	2008	46	2.17	3	na	3	0	99.91	3	93.99
Botticelli et al.	2008	21	5	0	0	0	0	105	0	100
Siegenthaler et al.	2007	29	1	0	0	0	0	29	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Ferrara et al.	2006	33	2.34	2	na	2	0	77.25	2.59	94.82
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Summary estimate (95%	% CI)*	424	1.80	6	0	6	1	675.28	0.81 (0.29–2.29)	98.4 (95.5–99.4)
Periodontal and non-pe	eriodontal rea	isons								
Gocen-Rohlig et al.	2010	20	2.33	0	0	0	0	46.67	0	100
Prosper et al.	2010	120	5	4	1	3	0	580.46	0.69	98.62
Mijiritsky et al.	2009	24	3.27	1	na	1	0	78.5	1.27	97.45
De Rouck et al.	2009	49	1.07	3	2	1	0	52.33	5.73	88.54
Crespi et al.	2009	64	2	0	na	0	0	128	0	100
Kan et al.	2009	20	2.15	0	na	0	0	43	0	100
Del Fabbro et al.	2009	61	1.79	1	1	0	0	107.67	0.93	98.14
Kahnberg	2009	40	2	0	0	0	0	80	0	100
Crespi et al.	2008	40	2	0	0	0	0	80	0	100
De Rouck et al.	2008	30	1	1	na	1	0	29.08	3.44	93.12
Cornelini et al.	2008	34	1	0	na	0	0	34	0	100
Covani et al.	2007	10	1	0	0	0	0	10	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Cangini and Cornelini	2005	32	1	0	0	0	0	32	0	100
Cornelini et al.	2005	22	1	0	na	0	0	22	0	100
Norton	2004	16	1.61	0	na	0	0	25.75	0	100
Covani et al.	2004	163	4	5	2	3	0	636.5	0.79	98.43
Bianchi et al.	2004	115	5.38	0	0	0	1	618.5	0	100
Becker et al.	1998	134	3.61	9	2	7	0	483.17	1.86	96.27
Lang et al.	1994	28	3.28	0	0	0	0	91.93	0	100
Becker et al.	1994	49	1.51	3	3	0	0	74.06	4.05	91.90
Summary estimate (95%	% CI)*	1094	2.24	27	11	16	1	3276.61	0.92 (0.09–9.15)	98.2 (83.3–99.8)

^{*}Based on random-effects *Poisson* regression, test for heterogeneity P = 0.01.

the extraction socket, 4.2% or 7 of 163 implants showed crestal bone contact apical to the first thread, among which six implants had bone contact between the first and second thread, and one implant presented bone loss up to the level of the third thread. Unfortunately, this study did not provide information on BOP.

Based on the limited studies and the inconsistent ways of assessing the soft tissues, it was difficult to estimate the prevalence of peri-implantitis.

Technical complications

Only three studies assessed technical complications that occurred in an observation period \geq 3 years. Lang et al. (1994) reported that "the reconstructions were complicationfree." Prosper et al. (2010) stated that "no pain or mobility of the definitive prosthesis

was registered" in the 5-year follow-up period.

Covani et al. (2004b) provided more detailed information concerning prosthetic complications. In this 4-year study, no fractures of abutments and/or prosthetic screws were documented. No prostheses needed to be replaced. The only prosthetic complication that occurred (9.8% of implants) was the loosening of the abutment screw.

				No of failure					Ertimotod	
		-	Mean	at 1 year					failure rate	Estimated
Study	Year of publication	Total no. of implants	follow-up time (year)	after implant placement	Before loading	After Ioading	Lost to follow-up	Total implant exposure time	(per 100 implant years)	survival after 2 year (%)
Anterior										
Tortamano et al.	2010	12	1.5	0	0	0	0	18	0	100
Gocen-Rohlig et al.	2010	10	2.33	0 0	0 0	0 0	0 0	23.33	0 0	100
Vicespilet al.	2010	c 0C	ک 15			5 0		05 C A	5 6	001
Nail et al. Del Fabbro et al	2005	210	02 T					37.63		001
Canullo et al	6002	- 9	80.6		o eu			12.5	00	100
Calvo-Guirado et al.	2009	46		, o	0	, c	, 	46	0 0	100
Miliritsky et al.	2009	20	3.57	0 0	na	0 0	- 0	71.42	0 0	100
Botticelli et al.	2008	ц Б	ъ Г	0	0	0	0	25	0	100
De Rouck et al.	2008	21	<i>~</i>	-	na	-	0	20.08	4.98	90.04
Cornelini et al.	2008	13	-	0	na	0	0	13	0	100
Crespi et al.	2008	24	2	0	0	0	0	48	0	100
Siegenthaler et al.	2007	14	-	0	0	0	0	14	0	100
Covani et al.	2007	5	-	0	0	0	0	5	0	100
Kan et al.	2007	23	-	0	na	0	0	23	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Ferrara et al.	2006	26	2.38	2	na	2	0	61.83	3.23	93.53
Cangini and Cornelini	2005	15	-	0	0	0	0	15	0	100
Cornelini et al.	2005	6	-	0	na	0	0	6	0	100
Vanden Bogaerde et al.	2005	19	1.5	0	0	0	0	28.5	0	100
Norton	2004	15	1.58	0	na	0	0	23.75	0	100
Covani et al.	2004	82	4	2	-	-	0	321.83	0.62	98.76
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Goldstein et al.	2002	16	2.77	0	0	0	0	44.33	0	100
Summary estimate (95% Cl,	*(486	1.92	5	-	4	-	1010.92	0.45 (0.15–1.36)	99.1 (97.3–99.7)
Posterior										
Prosper et al.	2010	120	LO I	4	- 1	m	0	580.46	0.69	98.62
Gocen-Rohlig et al.	2010	10	2.33	0	0 0	0	0	23.33	0	100
Crespi et al.	2010	5 2	2	0,	0,	0 0	0 0	30	0	100
Del Fabbro et al.	6007	40	1./9	- 0	- (0 0	0 0	70.04	1.43	97.14
	6002	0 5	2.U8	- c				55.55 7 00	0,1	25 15
Calvo-Guirado et al	2003	ל ד ל					- c	13 33	7 5	/1./0 85
Siciliano et al	2009	ς μ		- 0		- 0	. c	15.0		100
De Rouck et al	2008	<u>,</u> σ		0 0	, eu	, c	, c	<u>ר</u> ס	0 0	100
Cafiero et al.	2008	82		0 0	0	0 0	0 0	82	0 0	100
Cornelini et al.	2008	21	-	0	na	0	0	21	0	100
Fugazzotto	2008	341	2.72	2	-	-	0	929.06	0.22	99.57
Crespi et al.	2008	16	2	0	0	0	0	32	0	100
Botticelli et al.	2008	16	ß	0	0	0	0	80	0	100
Siegenthaler et al.	2007	15 1		0 0	0 0	0 0	0 0	- 1 5 -	0 0	100
Covani et al.	7005	7 D		5 6		5 0	5 0	ט 1 איז	5 0	001
rerrara et al.	2006	- F	7.20					24.01	5 6	00
Cornelini et al	2005									001
Vanden Bogaerde et al	2005	<u>,</u>	 -					46 5	0 0	100
Covani et al	2002	- 6	<u>5</u>	, w) -	<u>م</u> د		314.67	0.95	98.09
Norton	2004	; -	- 2	n 0	na	10	0 0	2	0	100
Fugazzotto et al.	2002	46	1.40	0	0	0	0	64.33	0	100
Goldstein et al.	2002	31	3.68	0	0	0	0	114.17	0	100
Summary estimate (95% Cl	*(967	2.02	12	4	8	-	2532.72	0.54 (0.04–7.05)	98.9 (86.8–99.9)
*Based on random-effects /	Poisson regression	n, test for heteroge	eneity $P = 0.0488$.							

Table 9. Comparison of annual failure rates and survival of implants inserted in extraction sockets - maxilla vs. mandible

Study	Year of	Total no. of implants	Mean follow-up time (year)	No. of failure at 1 year after implant placement	Before	After	Lost to	Total implant exposure time	Estimated failure rate (per 100 implant vears)	Estimated survival after 2 year (%)
Maxilla	P					j			J,	
Tortamano et al	2010	12	15	0	n 2	0	0	10	0	100
Calva Guirada at al	2010	12	1.5	1	na	1	1	10	1.60	100
Calvo-Guirado et al.	2009	40	1 07	ו כ	11d 2	1	0	59.55	1.09 E 72	90.05 00 E /
Crospi et al	2009	49	1.07	5	2	0	0	52.55 90	5.75	00.54
Crespi et al.	2009	40	2	0	na	0	0	80 42	0	100
Nan et al.	2009	20	2.15	0	na	0	0	43	0	100
	2009	30	1.79	0	0	0	0	23.75	0	100
Canullo et al.	2009	22	2.08	0	na	1	0	45.85	1 77	100
Nijiritský et al.	2009	24	5.27		na	0	0	/8.5	1.27	97.45
Botticelli et al.	2008	16	5	0	0	0	0	80	0	100
Crespi et al.	2008	40	2	0	0	0	0	80	0	100
Ribeiro et al.	2008	46	2.17	3	na	3	0	99.91	3	93.99
Lops et al.	2008	32	1.15	0	0	1	0	30.92	0	100
De Rouck et al.	2008	30	1		na	1	0	29.08	3.44	93.12
Catlero et al.	2008	21	1	0	0	0	0	21	0	100
Cornelini et al.	2008	27	1	0	na	0	0	27	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Siegenthaler et al.	2007	23	1	0	0	0	0	23	0	100
Covani et al.	2007	/	1	0	0	0	0	/	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Ferrara et al.	2006	33	2.34	2	na	2	0	77.25	2.59	94.82
Cangini and Cornelini	2005	15	1	0	0	0	0	15	0	100
Cornelini et al.	2005	19	1	0	na	0	0	19	0	100
Vanden Bogaerde et al.	2005	39	1.5	0	0	0	0	58.5	0	100
Tsirlis	2005	28	2	0	0	0	0	56	0	100
Norton	2004	16	1.61	0	na	0	0	25.75	0	100
Covani et al.	2004	95	4	2	1	1	0	373.75	0.54	98.93
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Fugazzotto et al.	2002	46	1.40	0	0	0	0	64.33	0	100
Goldstein et al.	2002	47	3.37	0	0	0	0	158.5	0	100
Summary estimate (95% CI) [*]		933	1.79	13	3	10	1	1805.46	0.73 (0.06–8.28)	98.6 (84.7–99.9)
Mandible										
Gocen-Rohlig et al.	2010	20	2.33	0	0	0	0	46.67	0	100
Prosper et al.	2010	120	5	4	1	3	0	580.46	0.69	98.62
Crespi et al.	2009	24	2	0	na	0	0	48	0	100
Del Fabbro et al.	2009	31	1.79	1	1	0	0	53.92	1.85	96.29
Lops et al.	2008	14	1.15	0	0	0	0	16.15	0	100
Cafiero et al.	2008	61	1	0	0	0	0	61	0	100
Cornelini et al.	2008	7	1	0	na	0	0	7	0	100
Fugazzotto	2008	341	2.72	2	1	1	0	929.06	0.22	99.57
Botticelli et al.	2008	5	5	0	0	0	0	25	0	100
Siegenthaler et al.	2007	6	1	0	0	0	0	6	0	100
Covani et al.	2007	3	1	0	0	0	0	3	0	100
Cangini and Cornelini	2005	17	1	0	0	0	0	17	0	100
Cornelini et al.	2005	3	1	0	na	0	0	3	0	100
Vanden Bogaerde et al.	2005	11	1.5	0	0	0	0	16.5	0	100
Covani et al.	2004	68	4	3	1	2	0	262.75	0.38	99.24
Summary estimate (95% Cl) [*]		731	2.10	10	4	6	0	2075.50	0.50 (0.17–1.52)	99 (97.0–99.7)
*Based on random-effect	s Poisson rear	ession test	for heteroger	peity $P = 0.05$						

In conclusion, there was insufficient data provided by the included papers to quantify technical/prosthetic complications.

Aesthetic outcomes

Only two studies (Bianchi & Sanfilippo 2004; Botticelli et al. 2008) evaluated the aesthetic results. In one study (Bianchi & Sanfilippo 2004), the buccal level of mucosal margin at prosthetic crown was compared with the level of the buccal gingival margin of the mesial and distal adjacent closest tooth crown. The threshold for the acceptable discrepancy was set at 1 mm. The data demonstrated complete success for the first 3 years, in the group of patients who received immediate implant placement and simultaneous connective tissue grafting, while only 80% of patients who were solely treated with immediate implants were considered successful. In the following 6 years, a small increase in the number of patients (<5%) who presented discrepancies of >1 mm was observed in both groups. The study concluded that soft tissue levels were generally stable following immediate implant placement, when connective tissue grafting was performed.

In the second study (Botticelli et al. 2008), the position of the mucosal margin during a 5-year period was followed. The mucosal margin moved 0.3 mm coronally at the proximal aspects, while there was an overall reces-

Table 10. Comparison of annual failure rates and survival of implants inserted in extraction sockets - loading protocol (immediate vs. conventional)

Study	Year of publication	Total no. of implants	Mean follow-up time (year)	No. of failure at 1 year after implant placement	Before loading	After loading	Lost to follow-up	Total implant exposure time	Estimated failure rate (per 100 implant years)	Estimated survival after 2 year (%)
Immediate loading										
Tortamano et al.	2010	12	1.5	0	na	0	0	18	0	100
Prosper et al.	2010	60	5	2	na	2	0	290.16	0.69	98.62
Mijiritsky et al.	2009	24	3.27	1	na	1	0	78.5	1.27	97.45
Calvo-Guirado et al.	2009	61	1	1	na	1	1	59.33	1.69	96.63
De Rouck et al.	2009	24	1	1	na	1	0	23.08	4.33	91.34
Crespi et al.	2009	64	2	0	na	0	0	128	0	100
Kan et al.	2009	20	2.15	0	na	0	0	43	0	100
Canullo et al.	2009	22	2.08	0	na	0	0	45.83	0	100
De Rouck et al.	2008	30	1	1	na	1	0	29.08	3.44	93.12
Cornelini et al.	2008	34	1	0	na	0	0	34	0	100
Crespi et al.	2008	20	2	0	na	0	0	40	0	100
Ribeiro et al.	2008	46	2.17	3	na	3	0	99.91	3	93.99
Crespi et al.	2007	150	1.5	0	na	0	0	225	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Siegenthaler et al.	2007	7	1	0	na	0	0	7	0	100
Kan et al.	2007	23	1	0	na	0	0	23	0	100
Ferrara et al.	2006	33	2.34	2	na	2	0	77.25	2.59	94.82
Barone et al.	2006	18	1	1	na	1	0	17.08	5.86	88.29
Cornelini et al.	2005	22	1	0	na	0	0	22	0	100
Vanden Bogaerde et al.	2005	50	1.5	0	na	0	0	75	0	100
Tsirlis	2005	28	2	0	na	0	0	56	0	100
Norton	2004	16	1.61	0	na	0	0	25.75	0	100
Kan et al.	2003	35	1.39	0	na	0	0	48.71	0	100
Summary estimate		822	1.72	12	na	12	1	1488.69	0.89	98.2
(95% CI) [^]									(0.18–4.40)	(91.6–99.6)
Conventional loading										
Vidal et al.	2010	54	1	0	0	0	0	54	0	100
Gocen-Rohlig et al.	2010	20	2.33	0	0	0	0	46.67	0	100
Crespi et al.	2010	30	2	0	0	0	0	60	0	100
Prosper et al.	2010	60	3	2	1	1	0	290.29	0.69	98.62
Del Fabbro et al.	2009	61	1.79	1	1	0	0	107.67	0.93	98.14
Kahnberg	2009	40	2	0	0	0	0	80	0	100
Cordaro et al.	2009	30	1.5	1	0	1	0	43.73	2.29	95.43
De Rouck et al.	2009	25	1.17	2	2	0	0	29.25	6.84	86.32
Siciliano et al.	2009	15	1	0	0	0	0	15	0	100
Botticelli et al.	2008	21	5	0	0	0	0	105	0	100
Lops et al.	2008	46	1.15	0	0	0	0	53.08	0	100
Catiero et al.	2008	82	1	0	0	0	0	82	0	100
Fugazzotto	2008	341	2.72	2	1	1	0	929.06	0.22	99.57
Crespi et al.	2008	20	2	0	0	0	0	40	0	100
Siegenthaler et al.	2007	22	1	0	0	0	0	22	0	100
Covani et al.	2007	10	1	0	0	0	0	10	0	100
Juodzbalys et al.	2007	14	2	0	0	0	0	28	0	100
Lineboom et al.	2006	25	1	2	2	0	0	24	8.33	83.33
Cangini and Cornelini	2005	32	1	0	0	0	0	32	0	100
Fugazzotto et al.	2002	46	1.40	0	0	0	0	64.33	0	100
Covani et al.	2004	163	4	5	2	3	0	636.5	0.79	98.43
Bianchi et al.	2004	115	5.38	0	0	0	1	618.5	0	100
Goldstein et al.	2002	4/	3.37	0	0	0	0	158.5	0	100
Huys Declare et al	2001	556	/	19	19	0	0	3/63./5	0.50	98.99
Becker et al.	1994	49	1.51	3	3	0	0	/4.06	4.05	91.90
Lang et al.	1994	28	3.28	0	0	0	0	91.93	0	100
Becker et al.	1998	134	3.61	9	2	/	0	483.17	1.86	96.27
(95% CI)*		2086	2.34	46	33	13	1	/942.48	0.75 (0.40–1.40)	98.5 (97.2–99.2)
*Based on random-effects <i>Poisson</i> regression, test for heterogeneity $P < 0.01$.										

sion of 0.4 mm at the buccal and 0.5 mm at the lingual aspects, respectively. At the 5year examination, the marginal level of the mucosa was generally coronal to the finishing line of the restorations. However, 5 of 21 buccal sites (24%) exhibited soft tissue recession to the extent that the margins of the metal restorations were exposed.

In summary, 20–25% of the patients who were treated with implants immediately placed into extraction sockets suffered aesthetically from apical displacement of the mucosal margins, although soft tissue levels seemed to be stable in long term in the majority of patients. The paucity of existing studies with follow-up periods of ≥ 3 years reporting on aesthetic outcomes following immediate implant placement made it diffi-

cult to estimate the prevalence of aesthetic complications and to investigate factors that might affect aesthetic outcomes.

Soft tissue changes

Three studies (Kan et al. 2003; De Rouck et al. 2008a; De Rouck et al. 2009) provided data on the soft tissue level changes, at 3, 6 and 12 months following implant placement immediately into extraction sockets and immediate provisional restoration, in relation to the pre-operative status in the anterior maxilla.

While the first two studies (Kan et al. 2003; De Rouck et al. 2008a) were prospective studies examining implants placed and restored immediately after tooth extraction, the latter study (De Rouck et al. 2009) was an RCT comparing the impact of immediate and delayed restorations on the soft and hard tissues following implantation. To obtain weighted means from comparable data, information from the immediate restoration group was extracted. The chi-squared tests showed that data from these three studies concerning changes in mesial and distal papilla heights and buccal mucosal levels at any examination time were homogeneous.

Alterations in the mid-buccal soft tissue level, and mesial and distal papilla heights were evaluated by measuring the distance from a reference line in one study (Kan et al. 2003). Acrylic stents, indented with mesial, buccal and distal grooves, were utilized to assess the papilla and mid-facial mucosal level in the two other studies (De Rouck et al. 2008a, 2009).

Fig. 3 shows the alterations in the mesial and distal papillae heights, and the mid-facial mucosal level in the first year following immediate implant placement and immediate restoration. Most of the soft tissue changes occurred in the first 3 months. Mesial and distal papillae shrank by 0.41 ± 0.32 mm (Q = 0.15, P = 0.93 and 0.34 \pm 0.36 mm (Q = 0.08, P = 0.96), respectively, while the buccal mucosal level was displaced apically by $0.43 \pm 0.38 \text{ mm} (Q = 0.01, P = 0.995), \text{ when}$ compared to the pre-surgical level. Soft tissues became stable after 6 months. At the end of the first year, 0.49 ± 0.31 mm (Q = 0.03, P = 0.99, 0.36 ± 0.33 mm (Q = 0.01, P = 0.99) and 0.51 ± 0.38 mm (Q = 0.02, P = 0.99) had been lost at mesial papilla, distal papilla and mid-facial mucosa, respectively.

Soft tissue alterations (mesial, distal and buccal), in the case of conventional loading, were evaluated by Cordaro et al. (2009), where 3 months of either submerged or nonsubmerged healing was allowed before



Fig. 3. Soft tissue changes in the first year after immediate implant placement and immediate restoration.

implants were loaded with provisional restorations. The measurements were taken at the time of provisional installation, and repeated at 3 and 15 months. This study used the incisal margin of adjacent teeth as the reference. As no statistically significant differences were found at any sites between the submerged and non-submerged implant groups, weighted means at each examination visit were calculated for the three sites. When compared to the pre-surgical soft tissue levels, the greatest loss was recorded at the time of provisional restoration (mesial papilla: -0.95 mm, distal papilla: -0.87 mm, buccal: -0.79 mm), after which, little changes had taken place (Fig. 4).

When soft tissue alterations upon immediate restoration were compared to those after delayed restoration, mean papilla shrinkage was about twice as high in the delayed restoration group (DRG) as the immediate restora-

tion group (IRG) at 3 months after provisional restorations (De Rouck et al. 2009). However, in the following 9 months, papillae in the DRG showed tendency to fill the proximal spaces, and the differences between the groups became smaller. On the other hand, mid-facial soft tissue loss showed little or no variation over time in both groups. The apical displacement of the buccal mucosal level was always about 2-3 times the magnitude in the DRG compared to the IRG during the 1-year observation period. It was concluded that this difference favoured immediate restoration (De Rouck et al. 2009) (Fig. 5a-c).

Papilla fill

Five studies provided information on the papilla fill after immediate implant placement. Implants in three studies (Cornelini et al. 2005; Kan et al. 2007a; Cornelini et al.



Fig. 4. Soft tissue changes after immediate implant placement and conventional loading [months after provisionals] (Cordaro et al. 2009): provisionals were installed 3 months after implant placement.

2008) were immediately restored, while implants in the other two studies (Lindeboom et al. 2006a; Juodzbalys & Wang 2007) were conventionally restored. All of them used the papilla index (Jemt 1997) to describe the fullness of the papillary fill.

The former three studies involved a total of 156 papillae. At the end of the observation period, 51% of the papillae achieved a score 2, i.e. the papilla was greater than half the height of the proximal space. The remaining

49% achieved a score 3, i.e. the papilla fills the entire proximal space. Fifty-one papillae were assessed after conventional loading. Forty-five percent belonged to score 2 and 55% achieved score 3.

Kan et al. (2007a) also investigated the change of distribution of papilla fill within the first year following immediate implant placement and immediate restoration. Of the 44 papillae examined, more than 90% scored 2 or 3 at every examination visit, and the



Fig. 5. Soft tissue changes – immediate restoration group (IRG) vs. delayed restoration group (DRG) (De Rouck et al. 2009). (a) Distal papilla. (b) Mesial papilla. (c) Buccal mucosal level.

number of papillae achieving score 3 continued to increase from implant placement and provisional insertion up until 6 months, after which papillae seemed to be stable (Fig. 6).

Pink Aesthetic Score (PES)

Only one study in the 46 included evaluated soft tissues using the PES (Juodzbalys & Wang 2007). Fourteen implants were placed in the region of upper incisors in 12 patients. GBR technique was employed. In case of soft tissue deficiency, connective tissue graft obtained from the palatal vault was used to cover the implant. Implants were firstly restored with provisional restorations at 6 month, then with definitive crowns at 12 month. One year after definitive crown cementation, the mean PES was 11.1. Incomplete mesial and distal papillae (64.3%), and alveolar process deficiency (42.9%) were common, and a minor discrepancy of buccal mucosal level of 1-2 mm was observed in 21.4% of cases.

Hard tissue change

Immediate implant placement and immediate loading

Generally, immediate implants in most studies experienced bone loss. The 1-year studies showed that the loss was less than 1 mm (range: gain 1 mm–loss 0.98 mm) in the first year, and longer-term studies demonstrated that after the first functioning year bone levels became stable.

De Rouck et al. (2008a, 2009) described the longitudinal radiographic marginal bony changes at 3, 6 and 12 months after immediate implant placement and immediate provisional restoration. The weighted means showed that from 3 to 12 months, there was a continuous loss of marginal bone from 0.51 ± 0.24 mm (Q = 0.05, P = 0.83) to 0.95 ± 0.35 mm (Q = 0.01, P = 0.93) at the mesial site, and from 0.52 ± 0.46 mm (Q =0.01, P = 0.91) to 0.79 ± 0.39 mm (Q =0.0001, P = 0.99) at the distal site. Half of the bone loss measured in the first year occurred in the first 3 months (Fig. 7).

Three studies (Calvo-Guirado et al. 2009; Crespi et al. 2009a; Canullo et al. 2009a) reported changes of marginal bone levels around immediately placed and immediately restored implants using platform-switching method. In the study of Calvo-Guirado et al. (2009), the mean bone loss after 1 year of function was 0.08 mm on the mesial surfaces and 0.09 mm on the distal surfaces. The small bony changes were in accordance with those reported in the RCT by Canullo et al.



Fig. 6. Change of distribution of papilla fill in the first year following immediate implant placement and immediate restoration. (Kan et al. 2007a,b)

(2009a), which showed that after about 2 years of loading, the platform-switching group experienced bone loss of 0.25 mm mesially and 0.36 mm distally; the bone loss was more significant in the platform-matching group, reaching 1.13 mm and 1.25 mm on mesial and distal surfaces, respectively. On the contrary, in the study by Crespi et al. (2009a), no significant differences in the bony changes between the two groups were found. The bone loss ranged 0.73–0.84 mm at the 1year follow-up and 0.68–0.80 mm at the end of second year.

Immediate implant placement and conventional loading

Authors used different baselines to measure bony alterations around implants loaded conventionally. Some used the bone level at the time of implant placement as the baseline, while the others chose the bone level at the time of implant loading to be the baseline.

For those with baseline at implant placement, bone loss of 1.01, 1.16 and 0.05 mm after 5 years, 21 months and 15 months of loading were reported, respectively (Crespi et al. 2008; Cordaro et al. 2009; Prosper et al. 2010).

Three studies used the bone level at the time of implant loading as the baseline (Juodzbalys & Wang 2007; Botticelli et al. 2008; Gokcen-Rohlig et al. 2010). Juodzbalys & Wang (2007) concluded that there was 1.16 mm bone loss 1 year after prosthetic restoration. Also, Gokcen-Rohlig et al. (2010) revealed bone loss of 0.72 mm and 1.36 mm at 1-year and 2-year follow-up, respectively. It was reported that two of four implants in one patient, which were used to support a fixed complete denture, experienced suppuration on probing; however, the suppuration resolved following local debridement and institution of strict oral hygiene practices. It was unclear if the substantial bone loss observed during the second year of function was due to biological complications or overloading. On the other hand, Botticelli et al. (2008) reported a mean bone gain of 0.2 mm after 5 years of loading (range: 0.22 mm loss to 0.41 mm gain). The minimal bony changes were attributed to the carefully supervised oral hygiene programme throughout the



Fig. 7. Hard tissue changes in the first year following immediate implant placement and immediate restoration.

whole observation period, with low plaque (11–17%) and bleeding (15–20%) scores at all follow-up visits.

Discussion

This systematic review showed that implants placed immediately in fresh extraction sockets yielded a low annual failure rate of 0.82% (95% CI: 0.48-1.39%) translating to a 2-year survival rate of 98.4%. It should be noted that this systematic review assumed a constant annual failure rate throughout the follow-up time after implant placement. With the knowledge that most implant failures occur in the first year, and that the current review has an annual failure rate based on studies with a mean observation period of 2 years, the rate derived should not be extrapolated to longer follow-up times. Therefore, studies with a mean follow-up period of 3 years or longer were analysed separately, reaching the 4-year implant survival rate of 97.5%. This percentage is comparable to the 96.8% 5-year survival rate of implants supporting single crowns reported in a previous systematic review (Jung et al. 2008).

In reference to the Third ITI Consensus Conference (Hämmerle et al. 2004), placement of implants is categorized by the healing timing following extraction as Type 1 immediate (within 24 h of extraction), Type 2 early (4–8 weeks after extraction), Type 3 early-delayed (12–16 weeks after extraction) and Type 4 late (more than 6 months). This classification is based on the time that elapsed after tooth extraction approximating the soft and hard tissue characteristics of healing sockets according to the morphologic, dimensional and histologic changes. In this systematic review, the survival rate of Type 1 placements was subject to evaluation.

However, a number of factors may affect the outcomes of procedures other than the timing of implant placement alone. Upon the timing of implantation, the type of the bone, the location and dimension of the edentulous area and the history of oral diseases are influential and should be considered in the assessment. On the other hand, the surgical protocol involved many steps influencing the outcomes as well. The approach chosen to perform the implant placement, such as with or without flap access, the selection of the implant type, the decision for the necessity of regenerative procedures and the selection of regeneration materials, the determination of the timing may all influence the outcomes of such procedures. Among the 46 studies in this systematic review, as many variables as possible were addressed. Nonetheless, five variables were discretely analysed.

Antibiotics

Infection should ideally be prevented after implant surgery. One of the proposed methods to minimize infection is the prescription of antibiotics to subjects undergoing implant surgery. The choice of antibiotics should be that it covers a reasonable bacterial spectrum to limit potential pathogens from colonizing in the vicinity of the surgical sites. When comparing subjects who had received a single dose pre-operatively, 5-7 days post-operatively, and a single dose pre-plus 5-7 days post-operative course of antibiotics, the estimated annual failure rates were 1.87%, 0.51% and 0.75% respectively. The annual implant failure rate in patients who were only given the single-dose of antibiotics pre-operatively was statistically significantly greater. This demonstrated that secondary to the prescription of an effective antibiotic, the duration of usage might be of importance. A single dose of antibiotics prior to surgery did not sustain the suppression of bacterial levels below the critical threshold throughout the healing period, but provision of antibiotics for 5-7 days after surgery may have helped to prevent post-operative infection, and hence, contribute to higher implant survival rates. However, these findings should be interpreted with caution, as the number of implants included in the single-dose pre-operative antibiotics group was substantially fewer.

Reasons for extraction

Implant sites with a history of periodontal disease may yield decreased survival rates. Many studies showed significantly more biological complications (Karoussis et al. 2003), greater peri-implant marginal bone loss (Mengel et al. 2007; De Boever et al. 2009), and increased implant failure rates (Hardt et al. 2002) in periodontitis susceptible subjects than periodontitis non-susceptible subjects. Furthermore, a recent review indicated that subjects with a history of periodontitis might be at greater risk for peri-implant infections (Renvert & Persson 2009). So, the comparison between implants placed into extraction sockets for non-periodontally related and periodontally related reasons was attempted in the present review. Unfortunately, not a single prospective study reported on a group of subjects with teeth extracted solely due to periodontal disease. Therefore, the comparison between implants placed in extraction sockets for non-periodontally related reasons, vs. mixed periodontal and non-periodontal reasons, was attempted. The two examined groups yielded comparable implant survival rates, although that of the non-periodontal group was slightly higher.

Site (maxilla/mandible)

Primary stability is of paramount importance for implant survival. Secondary to the dimensions of the extraction socket, the relative proportion of the load-bearing lamellar bone vs. cancellous bone also determines primary stability. As the mandible is comprised of a larger proportion of lamellar bone than in the maxilla, it is speculated that implant survival rates are correspondingly more favourable in the mandible. Accordingly, the 933 implants placed in the maxilla had an estimated annual failure rate of 0.73% compared to the 731 implants housed in the mandible, where the annual failure rate was 0.50%. This difference, however, was not statistically significant. One possible explanation could be that most of the studies adhered to a strict surgical protocol, where immediate implants were installed with a minimal insertion torque, and 3-4 mm apical bony engagement was ensured. Therefore, primary stability was achieved for all of the implants.

Site (anterior/posterior)

Due to the wider socket dimensions in multi-rooted tooth sites, it was speculated that there is a decreased amount of implant surface in direct contact with the adjacent bone walls. This might impede the achievement of optimal primary stability. However, as evidenced in this review, the difference in survival rates between implants placed in anterior single-rooted and posterior multirooted sockets was negligible. This could again, be attributed to the surgical protocol, where minimal insertion torque and engagement of bone apical to the socket was propagated. Moreover, usage of different diameter implants matched to the various socket dimensions could have contributed to the observed minimal difference.

Loading

Corroborating the findings by Gallucci et al. (2009), the utilized loading protocols did not result in any significant difference with regard to survival rates. Immediately loaded and conventionally loaded implants had reported implant survival rates of 98.2% and 98.5%, respectively after a 2-year observation period. In most of the studies reporting on immediate loading, the inserted restorations were free of contacts in centric occlusion and

during excursive movements; utilizing such an occlusal scheme, micromovements of implants were certainly limited. Therefore, it is not surprising to note that the differences in survival rates between the two groups with varying loading protocols were not significant.

Success

A successful treatment should be the treatment with absence of any biological, technical and aesthetic complications. Since all these complications take time to develop, this systematic review assessed a total of nine studies with the mean follow-up time of 3 years or more for the estimation of the success rates of implant-related therapy.

Biological complications

Peri-implant mucositis and peri-implantitis have been shown to be prevalent. A systematic review (Zitzmann & Berglundh 2008) concluded that peri-implant mucositis occurred in approximately 80% of the subjects and in 50% of the implants; while periimplantitis was found in 28% and \geq 56% of subjects and in up to 43% of implant sites.

Diagnosis of peri-implant diseases required assessment of the presence or absence of bleeding on probing (BOP) in the peri-implant soft tissues, and changes in the level of crestal bone (Lang & Berglundh 2011). Three of the nine studies reported on BOP. In one study (Bianchi & Sanfilippo 2004), 31% of the implants showed signs of peri-implant mucositis.

Seven studies described hard tissue conditions. An unusual amount of marginal bone loss was seen in 4.3% of implants in the study of Covani et al. (2004a,b). However, in the absence of the report on BOP, it was difficult to estimate the prevalence of peri-implantitis.

Technical complications

Three studies assessed technical complications, among which two were free from this type of complication and one study (Covani et al. 2004b) had loosening of abutment screws occurring in 9.8% of implants supporting single crowns during the 4 years of function. This finding is comparable to that reported in the systematic review on implant-supported single crowns (Jung et al. 2008), which demonstrated a cumulative incidence of screw or abutment loosening of 12.7% in 5 years. The technical complications in implant-supported reconstructions are generally three times as high as those in tooth-supported reconstructions (Pjetursson et al. 2007). This may be explained by the ankylotic union of implants and the bone, and the lacking of a periodontal ligament around implants, subsequently resulted in the increased threshold to mechanorecptive responses.

Aesthetic complications

Although not included in previous descriptions of implant success criteria (Albrektsson et al. 1986), the aesthetic aspect of implant-supported restorations has attracted more attention in recent years. Despite the high survival rate of immediate implants, which has been addressed in this systematic review, soft tissue alterations, especially the buccal marginal mucosal recession, appeared to be inevitable. About 20% of patients in the two included studies with follow-up time >3 years (Bianchi & Sanfilippo 2004; Botticelli et al. 2008) suffered from restorations with limited aesthetic outcomes due to buccal soft tissue recession. It was in accordance with a recent follow-up study by Kan et al. (2011), which stated that while the mean aesthetic satisfaction rating by patients was almost perfect (9.9 of 10) at the 1-year recall, 4 of 35 patients (11%) complained of unsatisfactory restorations caused by facial gingival recession after a longer period of observation (mean 4 years, range: 2-8.2 years). Three of these subjects agreed to undergo additional guided bone regeneration and connective tissue grafting surgeries to correct the problem.

When immediate implant placement is indicated, careful case selection and cautious treatment planning are essential to minimize aesthetic complications. Buccal soft tissue recession has been shown to be closely related to the thin tissue biotype (Kan et al. 2011) and buccally positioned implants (Chen et al. 2007). In case of thin tissue biotype, connective tissue grafting may have to be performed (Bianchi & Sanfilippo 2004).

In recent years, several indices have been developed to provide guidance on objective and comprehensive assessment of aesthetic outcomes of an implant restoration. They include the Pink Esthetic Score (Fürhauser et al. 2005), the Implant Crown Aesthetic Index (Meijer et al. 2005) and the modified PES/White Esthetic Score (WES) (Belser et al. 2009). However, none of the long-term studies in this systematic review evaluated the aesthetic outcomes using any of these indices. In the future, more routine utilization of these indices is recommended for aesthetic monitoring.

Tissue changes

Soft tissue change

It was observed in three studies (Kan et al. 2003; De Rouck et al. 2008a, 2009) that the bulk of the soft tissue changes occurred during the first 3 months of healing after immediate implant placement and immediate restoration. Change of a smaller magnitude was exhibited in the following 3 months but stabilized after the first 6 months. At the end of the first year, the weighted mean loss at mesial papilla, distal papilla and mid-facial mucosa were 0.49 mm, 0.36 mm and 0.51 mm, respectively.

In a recent publication, Kan et al. (2011) followed up the same patient population as the study published in 2003 for 2-8.2 years (mean 4 years) and reported the soft tissue changes beyond the first year evaluation. When compared to the pre-surgical status, mesial and distal papillae lost height of 0.53 mm and 0.39 mm at first year followup; and lost 0.22 mm and 0.21 mm at the last examination appointment. The significantly smaller loss in papilla height over time demonstrated that papillae might have the capacity of continuous regrowing following implant restoration. It is also important to note that gingival biotype did not significantly influence the papilla level changes. Both biotypes showed loss in height of 0.21 mm at last examination visit. The papillary height might be more likely to be affected by factors such as proximal bone level of neighbouring teeth and the distance between the implant and adjacent tooth.

On the other hand, significantly more recession was reported at the facial mucosa at the last examination visit than the firstyear follow-up (-1.13 mm vs. -0.55 mm), and significantly more apical displacement occurred in patients with thin gingival biotype than those with thick gingival biotype (-1.50 mm vs. -0.56 mm). In a closer look at the mean facial gingival level changes at the 1-year and final (mean of 4 years) examinations, the subjects with a thin biotype had a greater 1-year mean recession $(-0.75 \pm 0.59 \text{ mm})$ than the final recession observed in subjects who carry a thick biotype at $(-0.56 \pm 0.46 \text{ mm})$. Needless to say, the subjects with a thin biotype had the most recession at the final observation $(-1.5 \pm 0.88 \text{ mm})$. Therefore, it is speculated that these limited aesthetic results were mainly witnessed in subjects with a thin biotype and only gradually manifested at a later life of the immediate implant supported restoration caused by progressive facial gingival recession.

The above results showed that although the greatest changes in soft tissues took place in the first 6 months following immediate implant placement and immediate restoration, soft tissue remodelling might continue over the years. While mesial and distal papillae had tendencies to gain height, buccal mucosal recession might get more pronounced over time.

Apical displacement of the mucosa was also inevitable in the case of delayed loading. Cordaro et al. (2009) concluded that the greatest loss was recorded at the time of provisional restoration, which was carried out 3 months after implant installation (mesial papilla: -0.95 mm, distal papilla: -0.87 mm, buccal: -0.79 mm). In the following 9 months of the study, little changes had taken place (Fig. 3).

One randomized clinical trial (De Rouck et al. 2009) compared longitudinal changes of papilla height and position of the buccal mucosa between immediately and conventionally restored implants after immediate implant placement. Papilla shrinkage and the apical displacement of buccal mucosa were of a lesser extent in immediate restored implant group at the 3-month follow-up. Nevertheless, at the 12-month re-examination, the two groups yielded comparable results when examining the change in papilla height. However, the differences in the position of the buccal mucosa persisted throughout the 12-month observation period. The authors thus concluded that immediate restoration of immediate implants might help limit buccal recession, but more randomized clinical trials of longer follow-up period are required before any definitive conclusions can be drawn on this issue.

In addition to the dimensional changes of soft tissues, the prevalence of mucosal recession should also be noted. Chen et al. (2007a, b) investigated this aspect among immediately placed delayed restored implants. In this prospective clinical study, 30 patients were randomly assigned to one of the three groups: (i) bone graft group (BG), (ii) bone graft and resorbable membrane group (BG +M), and (iii) non-grafted group (control). After 6 months of initial healing, 10 of 30 sites (33.3%) showed buccal marginal tissue recession (range: 1-3 mm), where three were from the BG group, four from the BG+M group and three from the control group. Five of the 10 patients with recession and two patients without recession then received connective tissue grafts to repair or prevent the recession. At the time of final crown installation at 8 months, 8 of 30 sites exhibited marginal tissue recession when compared to contralateral teeth and thus suboptimal aesthetic results, among which three sites had previously received connective tissue grafts.

The current evidence indicated that implants placed in extraction sockets were not able to prevent soft tissue loss, especially the buccal marginal tissue recession. The amount of soft tissue alterations, however, was determined by many factors. While potential benefits of immediate restoration of an implant required further investigation, some fundamental factors should not be overlooked to minimize recession. The prospective study mentioned (Chen et al. 2007) demonstrated a significant relationship between the frequency of recession and the bucco-lingual position of the implants. Six of the eight implants, which showed marginal tissue recession, were placed at or buccal to the reference line joining the buccal cervical margins of adjacent teeth. Similarly, a retrospective study (Chen et al. 2009) also noted that implants placed more buccally experienced greater recession than implants placed lingually (-6.9% vs. -2.6%).

This latter concept has also been confirmed with re-entry surgery at 4 months in a randomized controlled clinical trial of implants installed immediately into extraction sockets (Tomasi et al. 2010).

Furthermore, recession of >5% was more prevalent at sites with thin periodontal biotypes than at those with a thick biotype. Therefore, when immediate implant placement is indicated, careful pre-surgical examination of future implant sites and placement of implants in the prosthetically correct position should be carried out to achieve and maintain satisfactory aesthetic outcomes.

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Hard tissue change

As with the soft tissue changes, most of the marginal bone loss was found in the first 3 months following immediate implant placement and immediate restoration. At the end of the first year, the bone loss was generally less than 1 mm. One possible means to minimize the hard tissue changes, in shorttem, could be the use of platform-switching technique, where a wider-diameter implant is restored with a narrower-diameter abutment. In one randomized clinical controlled trial, significantly less mean bone resorption occurred adjacent to platform-switched abutment restorations than that found at sites using platform-matched abutments (Canullo et al. 2009a). However, in another RCT, no such differences were demonstrated (Crespi et al. 2009a). Hence, more clinical trials are required to confirm the possible benefits of the platform-switching technique.

In the long run, good oral hygiene is a prerequisite for maintaining bone levels. With low plaque and mucositis levels, bone levels even improved (mean gain of 0.2 mm) after 5 years of implant functioning (Botticelli et al. 2008).

Conclusion

- The estimated annual failure rate of implants placed in extraction sockets was 0.82% (95% CI: 0.48–1.39%) translating to a 2-year survival rate of 98.4% (97.3–99%).
- 2. The estimated annual implant failure rate was lower after a 5–7 days post-operative antibiotic course (0.51%) than a single dose of pre-operative antibiotics (1.87%) (P = 0.002).
- Scarce data concerning biological complications were available in long-term (≥3 years) studies. Future research

- should pay more attention to evaluate peri-implant tissues by periodontal probing and radiographs.
- 4. Technical complications were not commonly reported in studies with follow-up time of 3 years or more.
- 5. About 20% of patients who underwent immediate implant placement and delayed restorations suffered from suboptimal aesthetic outcomes due to buccal soft tissue recession in studies with observation period of 3 years or more.
- 6. It has been shown in a 1-year RCT that immediate restoration after immediate implant placement might help limit buccal mucosal recession, but more longterm RCTs are required to confirm this potential benefit. On the other hand, the influence of factors, such as gingival biotypes and bucco-lingual position of implants, on buccal soft tissue levels should not be overlooked.
- 7. To date the use of platform-switching technique to reduce marginal bone resorption is controversial and needs further investigation. However, good OH is still a pre-requisite for maintaining bone levels in the long run.

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List of excluded full text articles and the reason for exclusion

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- Gatti, C., Haefliger, W. & Chiapasco, M. (2000) Implant-retained mandibular overdentures with immediate loading: a prospective study of iti implants. The International Journal of Oral & Maxillofacial Implants 15: 383–388. Exclusion criteria: Sample size (subject) <10 in the immediate implant group.
- Glauser, R., Ree, A., Lundgren, A., Gottlow, J., Hammerle, C.H. & Scharer, P. (2001) Immediate occlusal loading of branemark implants applied in various jawbone regions: a prospective, 1-year clinical study. *Clinical Implant Dentistry & Related Research* 3: 204–213. *Exclusion criteria: unknown survival rate of immediate implants; no separate report of the survival rate of immediate implants from other types of implant placement.*

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- Glauser, R., Zembic, A., Ruhstaller, P. & Windisch, S. (2007) Five-year results of implants with an oxidized surface placed predominantly in soft quality bone and subjected to immediate occlusal loading. *Journal of Prosthetic Dentistry* 97: S59– 68. Exclusion criteria: not reporting on immediate implants.
- Gomez-Roman, G., Kruppenbacher, M., Weber, H. & Schulte, W. (2001) Immediate postextraction implant placement with root-analog stepped implants: surgical procedure and statistical outcome after 6 years. The International Journal of Oral & Maxillofacial Implants 16: 503-513. Exclusion criteria: unknown survival rate of implants in extraction sockets; no separate report of the survival rate of implants in extraction sockets from implants at explantation sites.
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- Guncu, G.N., Tozum, T.F., Guncu, M.B., Yamalik, N. & Tumer, C. (2009) A 12-month evaluation of nitrite oxide metabolism around immediate and conventionally loaded dental implants. *Implant* Dentistry 18: 27–37. Exclusion criteria: not reporting on immediate implants.
- Hall, J.A., Payne, A.G., Purton, D.G., Torr, B., Duncan, W.J. & De Silva, R.K. (2007) Immediately restored, single-tapered implants in the anterior maxilla: prosthodontic and aesthetic outcomes after 1 year. *Clinical Implant Dentistry* & *Related Research* 9: 34–45. *Exclusion criteria: not reporting on immediate implants.*
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 108: e19–25. Exclusion criteria: mean follow-up time < 1 year.

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- Hruska, A., Borelli, P., Bordanaro, A.C., Marzaduri, E. & Hruska, K.L. (2002) Immediate loading implants: a clinical report of 1301 implants. *Journal of Oral Implantology* 28: 200–209. *Exclusion criteria: not reporting on immediate implants.*
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