

Gerry M. Raghoobar, Henny J.A. Meijer, Wim Slot, James J.R. Huddleston Slater, Arjan Vissink

A systematic review of implant-supported overdentures in the edentulous maxilla, compared to the mandible: How many implants?



Gerry M. Raghoobar, DDS, MD, PhD

Professor, University of Groningen, University Medical Center Groningen, Department of Oral and Maxillofacial Surgery, Groningen, The Netherlands

Henny J.A. Meijer, DDS, PhD

Professor, University of Groningen, University Medical Center Groningen, Department of Oral and Maxillofacial Surgery, and Department of Oral Function and Prosthetic Dentistry, Groningen, The Netherlands

Wim Slot, DDS, PhD

Assistant Professor, University of Groningen, University Medical Center Groningen, Department of Oral Function and Prosthetic Dentistry, Groningen, The Netherlands

James J.R. Huddleston Slater, DDS, PhD

Assistant Professor, University of Groningen, University Medical Center Groningen, Department of Oral and Maxillofacial Surgery, Groningen, The Netherlands

Arjan Vissink, DDS, MD, PhD

Professor, University of Groningen, University Medical Center Groningen, Department of Oral and Maxillofacial Surgery, Groningen, The Netherlands

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Background and aim: There is now overwhelming evidence from systematic reviews that a two-implant overdenture is the first choice of treatment for the edentulous mandible. Conversely, consensus is lacking for implant-supported maxillary overdentures. Therefore, we systematically reviewed the treatment outcome of concepts used for implant-supported maxillary overdentures, focusing on the survival of implants, survival of maxillary overdentures and condition of the implant surrounding hard and soft tissues after a mean observation period of at least 1 year.

Material and methods: MEDLINE (1950 to December 2013), EMBASE (1966 to December 2013) and CENTRAL (1800 to December 2013) were searched to identify eligible studies. Two reviewers independently assessed the articles using specific study design-related quality assessment forms.

Results: Out of 195 primarily selected articles, 24 studies fulfilled the inclusion criteria. A meta-analysis showed an implant survival rate of 98.1% and overdenture survival of 99.5% per year in the case of ≥ 6 implants and a splinted (bar) anchorage. In the case of ≤ 4 implants and a splinted (bar) anchorage, implant survival rate and overdenture survival were 97.0% and 96.9% per year, respectively. In the case of ≤ 4 implants and a non-splinted anchorage (ball, locator, telescopic crown), implant survival rate and overdenture survival were 88.9% and 98.8% per year, respectively. The condition of the peri-implant tissues was not reported in most studies.

Conclusions: An implant-supported maxillary overdenture (all studies ≥ 4 implants) provided with a splinted anchorage is accompanied with a high implant and overdenture survival rate (both $>95\%$ per year), while there is an increased risk of implant loss when ≤ 4 implants with a non-splinted anchorage are used.

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■ Introduction

Edentulous patients often experience serious functional and psychosocial problems related to their conventional dentures because of an impaired load-bearing capacity^{1,2}. These problems include pain

during mastication, and insufficient stability and retention of the denture. Resolving such problems, particularly before the advent of implants, has been a challenge for both the prosthodontist and surgeon.

More than 20 years ago, van Steenberghe et al³ first reported on the possibility of using man-

Correspondence to:

Prof Dr G.M. Raghoobar
 Department of Oral and
 Maxillofacial Surgery,
 University Medical Center
 Groningen,
 P.O. Box 30.001,
 9700 RB Groningen,
 The Netherlands
 Tel: +31503613840
 Fax: +31503611136
 Email: g.m.raghoobar@
 umcg.nl

Table 1 Search strategy.

| | |
|--|---|
| #1 Search | "Denture, Overlay" [MeSH] |
| #2 Search | "Dental Prosthesis, Implant supported" [MeSH] |
| #3 Search | "Dental Implants" [MeSH] |
| #4 Search | "Dental Implantation, Endosseous" [MeSH] |
| #5 Search | "Mouth, Edentulous" [MeSH] |
| #6 Search | "Jaw, Edentulous" [MeSH] |
| #7 Search | "Maxilla" [MeSH] |
| #8 Search | #2 OR #3 OR #4 |
| #9 Search | #5 OR #6 |
| #10 Search | #1 AND #7 AND #8 AND #9 |
| Last run of data search: 31 December, 2013 | |

dibular overdentures supported by two implants to treat problems where usually conventional mandibular dentures would be used. Since then, mandibular overdentures have been extensively studied with respect to a number of implants, a variety of clinical items (including implant survival, health of peri-implant soft tissues and peri-implant bone loss) and patients' satisfaction⁴⁻¹⁰. For the vast majority of patients, an overdenture on two implants in the mandible is the first choice of treatment when complaining about the lack of stability in their mandibular denture¹¹⁻¹³. Underlining the McGill and York consensus statements, Thomason et al¹⁴ concluded that there is now overwhelming evidence to support the proposal that a two-implant overdenture should become the first choice of treatment for the edentulous mandible. The number of implants in the edentulous mandible for support of an overdenture are well studied^{15,16}.

Regarding implant-supported maxillary overdentures, consensus is lacking, but implant-supported maxillary overdentures have been shown as a favourable treatment option for patients with persistent complaints of retention and stability of their conventional maxillary denture¹. Next to sufficient retention and stability, proper phonetics, aesthetics and hygiene access can be achieved with implant-supported maxillary overdentures.

While two endosseous implants are generally considered to provide sufficient support to a mandibular overdenture, the number of implants needed to support a maxillary overdenture is still not set.

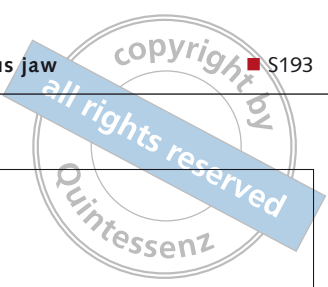
Currently, a variety of numbers of implants is applied to support the maxillary overdenture, as well as a variety of anchorage systems¹⁷. Sadowsky¹⁸ evaluated maxillary implant-supported overdentures with emphasis on the number of implants and anchorage design. He concluded that a number of 4 implants was the minimum to support a maxillary overdenture and recommended 6 implants in case of compromised bone. He could not detect a difference between the treatment outcome of splinted and non-splinted implants in the literature he assessed. Three years later, Slot et al¹⁹ showed in a meta-analysis that the survival of implants used to support a maxillary overdenture is high if concepts were used with at least 4 implants supplied with either a bar or ball anchorage. Finally, from the systematic review of Rocuzzo et al¹⁶, it can be concluded that the question of how many implants should support a maxillary overdenture is still open. Therefore, the aim of this systematic review was to assess the treatment outcome of concepts used for implant-supported maxillary overdentures focusing on survival of implants, survival of maxillary overdentures and the condition of surrounding hard and soft tissues after a mean observation period of at least 1 year.

■ Material and methods

■ Design of the study and search strategy

Although randomised controlled trials (RCTs) provide the highest evidence in comparing effectiveness of different therapies, relevant information is not exclusively provided by RCTs. Well-designed clinical trials and case series may also provide valuable information.

A search of the literature was conducted in the databases of MEDLINE (1950 to 31 December, 2013) (via PUBMED) and EMBASE (1966 to 31 December, 2013). The search was supplemented with a systematic search in the Cochrane Central Register of Controlled Trials' (CENTRAL) (1800-31 December, 2013). No language restriction was applied. The search strategy was a combination of MeSH terms (Table 1). The search was completed by checking the references of the relevant review articles and eligible studies.



Full-text documents were obtained for all articles meeting the inclusion criteria. Full text analysis was performed independently by two reviewers (GR, HM). Methodological quality was assessed independently by the reviewers using specific study design-related modified forms designed by the Dutch Cochrane Collaboration²⁰. In case of disagreement, a consensus was reached by discussion, if necessary in consultation with a third reviewer (AV). To ensure that datasets were unique, of the studies in which the same patients were analysed at different times, leading to different publications, the study with the longest follow-up was selected for definitive analysis.

The criteria for a paper to be included in the study selection were:

- detailed information on maxillary overdentures supported by root-form endosseous implants; in case of combined data for implant-supported maxillary and mandibular removable overdentures, extraction of data for the maxillary overdenture must be possible
- the treatment of the patients has to be initially planned for a maxillary overdenture
- at least five patients should be described in a paper
- the follow-up period for implants in maxilla should be at least 1 year
- study design: RCTs, clinical trials or case series; retrospective studies were excluded.

■ Outcome measures

The following outcome measures were assessed:

- survival of implants
- survival of overdentures
- condition of peri-implant hard and soft tissues.

■ Statistical analysis

For the meta-analysis, the statistical software package 'Meta-analysis' was used (Comprehensive Meta-analysis Version 2.2, Biostat, Englewood, NJ 2005). For the calculation of the overall effects for the included studies, weighted rates together with random effect models were used.

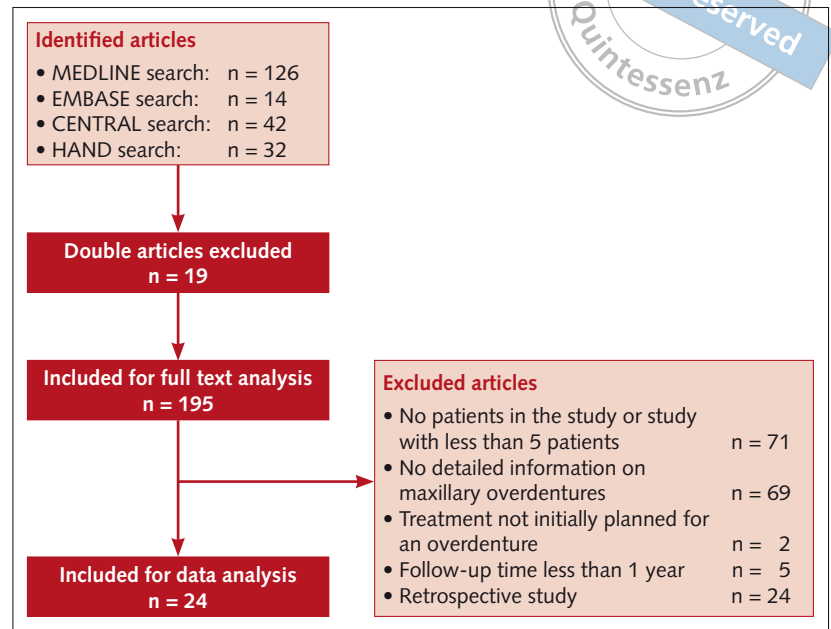
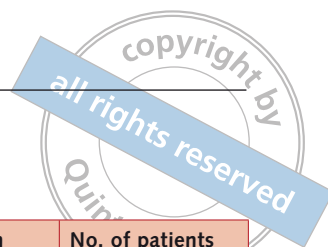


Fig 1 Algorithm of study selection procedure.

■ Results

■ Description of the studies

The MEDLINE search provided 126 hits, the EMBASE search 14 hits and the CENTRAL search 42 hits. Nineteen articles appeared to be duplicated. After scanning titles and abstracts, it was decided to select them all for evaluation as the full text article, because the abstracts did not always give a clear insight in the method of the study and the number of hits was reasonable to assess. This way no article was excluded beforehand. Reference-checking of relevant reviews and included studies revealed 32 additional articles to be screened. This approach resulted in 195 articles to be evaluated by full text analysis. Seventy-one articles were excluded because no patients at all or less than 5 patients were described. Another 69 articles were excluded because there was no detailed information available on maxillary overdentures as a separate treatment. Two articles were excluded because the treatment with implants was not initially planned for an overdenture. Five articles were excluded because the follow-up was less than 1 year. Finally, 24 articles were excluded because they were retrospective studies. The remaining 24 articles were scored (Fig 1).

**Table 2** General characteristics of included studies.

| Study | Year of publication | Study design | Follow-up in months | No. of patients in study |
|--------------------------------------|---------------------|---|---------------------|--------------------------|
| Zou et al ⁴⁴ | 2013 | Prospective | 36 | 30 |
| Slot et al ⁴¹ | 2014 | Prospective (Straumann group) | 12 | 25 |
| | | Prospective (Astra Tech group) | 12 | 25 |
| Slot et al ⁴³ | 2014 | Randomised Controlled Trial | 12 | 66 |
| Slot et al ⁴² | 2013 | Randomised Controlled Trial | 12 | 50 |
| El-Ghareeb et al ⁴⁰ | 2012 | Prospective | 14 | 6 |
| Van Assche et al ³⁹ | 2012 | Prospective | 24 | 12 |
| Katsoulis et al ³⁸ | 2011 | Prospective | 24 | 28 |
| Mangano et al ³⁷ | 2011 | Prospective | 60 | 38 |
| Akça et al ³⁶ | 2010 | Prospective | 59 | 11 |
| Pieri et al ³⁵ | 2009 | Prospective | 12 | 22 |
| Raghoobar et al ³⁴ | 2006 | Prospective | 22 | 8 |
| Raghoobar et al ³³ | 2005 | Prospective | 20 | 5 |
| Payne et al ³² | 2004 | Randomised Controlled Trial (Brånemark group) | 12 | 20 |
| | | Randomised Controlled Trial (Southern group) | 12 | 19 |
| Raghoobar et al ³¹ | 2003 | Prospective | 12 | 10 |
| Ferrigno et al ³⁰ | 2002 | Prospective | 120 | 35 |
| Zitzmann and Marinello ²⁹ | 2000 | Prospective | 12 | 10 |
| Zitzmann and Marinello ²⁸ | 2000 | Prospective | 27 | 10 |
| Bergendal and Engquist ²⁷ | 1998 | Randomised Controlled Trial (bar group) | 60 | 10 |
| | | Randomised Controlled Trial (ball group) | 50 | 8 |
| Naert et al ²⁶ | 1998 | Prospective | 48 | 13 |
| Watson et al ²² | 1997 | Prospective | 60 | 30 |
| Jemt et al ²¹ | 1996 | Prospective | 60 | 30 |
| Hutton et al ²⁵ | 1995 | Prospective | 36 | 30 |
| Jemt et al ²⁴ | 1994 | Prospective | 12 | 6 |
| Johns et al ²³ | 1992 | Prospective | 12 | 30 |

Two studies were suspected to present the same study population^{21,22}. Whether the same study population was used was not clearly stated in the manuscript and for this reason, it was doubtful. As these two studies deliver the same data for the meta-analysis, the data from the most recent manuscript was used for the meta-analysis²². Both studies were saved for the tables, however, as regards survival, the focus was on different evaluation items. The two disagreements that occurred were easily resolved in a consensus meeting.

General characteristics of the 24 included studies are depicted in Table 2²¹⁻⁴⁴. Authors of two articles^{35,38} responded to an email concerning queries regarding the different groups they mentioned in their article. In the latter study, patients with 5

implants were excluded³⁸. Four studies were randomised controlled trials (RCTs)^{27,32,42,43}. In the study of Payne et al³² two different implant systems were analysed and in the study of Bergendal and Engquist²⁷, the difference between a bar and a ball anchorage design was studied. In both studies, the patients that were included received 3 or less implants and a ball anchorage. Only one study was included regarding <4 implants provided with a bar suprastructure²². Slot et al^{42,43} reported on the 1-year treatment outcome of 4 and 6 bar-connected implants placed with or without pre-implant bone augmentation to support an overdenture in edentulous patients. There was no difference in implant loss between these groups. In a 3-year prospective study, Zou et al⁴⁴ evaluated the use of telescopic

crown, bar and locator attachments to support a removable 4 implant-supported maxillary overdenture. No significant differences were observed in the implant survival and success rates. Furthermore, they showed that the locator attachment system was accompanied with the best peri-implant hygiene, frequency of prosthodontic maintenance measures, costs and ease of denture preparation when compared to the telescopic crown and bar attachment systems. Slot et al⁴¹ also reported the results of a 1-year prospective case series in two groups of 25 patients on the treatment outcome of maxillary overdentures supported by 6 implants opposed by natural antagonistic teeth in the mandible. In the 25 patients in whom the implants were placed after augmentation, one implant was lost and in the 25 patients not needing pre-implant augmentation, three implants. The remaining 19 studies described prospectively analysed case series. The number of patients in the studies varied from five patients to 66 patients. The follow-up period varied from 12 to 120 months (Table 2).

Table 3 summarises the treatment procedures of the included studies. The number of implants placed to support the overdenture varied from 2 to 8 implants. Onlay block graft procedures and elevation of the floor of the maxillary sinus were carried out in some studies before insertion of the implants or together with the placement of the implants. Also, the placement of implants without bone graft procedures was described. The position of the implants, in relation to the availability of a bone volume sufficient to reliably insert endosseous implants, was often not well described. Furthermore, different implant systems were used (the majority were Brånemark and Straumann implants) as well as various anchorage systems. As regards anchorage systems, both splinted (bar) and non-splinted (ball, locator and telescopic crown) designs were used. With ≥ 6 implants, the anchorage design was splinted in all cases. With ≤ 4 implants both designs were used. In the majority of the studies, the kind of opposing dentition was not described; other studies described that there were all kinds of opposing dentition. Only in three RCTs^{32,42,43} was it mentioned that all patients had a 2-implant or 4-implant overdenture in the mandible.

Table 4 gives the outcomes of the studies included in this review. For the survival rates of implants and

overdentures, see the meta-analysis paragraph. The condition of the surrounding hard and soft tissues was mentioned in nine out of the 24 studies. In 13 studies, a change in mean marginal bone level was mentioned. When reported, a variety of outcome parameters were used, as measurements were done on either non-standardised rotational panoramic radiographs and intraoral radiographs, or on standardised intraoral radiographs. Loss of marginal bone varied from 0.22 mm in 12 months to 1.25 mm in 60 months. In 7 studies, the condition of the peri-implant mucosa was mentioned, but unfortunately a variety of indices was used to score this condition. In 8 studies, bleeding on probing was noted. Finally, in 7 studies probing depth was mentioned, varying from 3.2 mm to 4.8 mm.

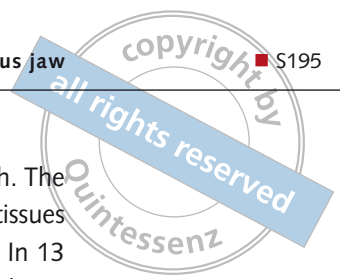
■ Meta-analysis

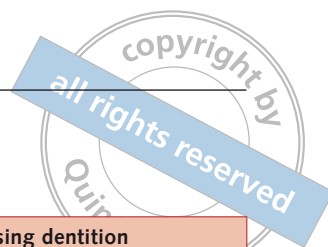
Due to the methodological diversity of the studies, only the number of implants, anchorage design, survival of implants and survival of the overdenture could be meaningfully combined in a meta-analysis. It was chosen to include ≥ 6 implants and ≤ 4 implants in the meta-analysis to have a clear distinction between these two groups.

Figs 2, 3 and 4 depict the results of the weighted meta-analysis, expressed as event rates per year. Event rates were used to describe failures and were calculated by the ratio of the number of failures or complications (e.g. events) to the total exposure time of the construction. The exposure time was the time the implants or the overdenture was followed. Distinct event rates were calculated for both implants and dentures. In case of an implant failure or dentures that were lost during the observation time, the time to the event was used for the analysis. The survival rate (SR) is the complement of the event rate (ER), and was calculated as $SR = 1 - ER$.

■ Survival of implants

Implant survival was defined as the percentage of implants initially placed that was still present at follow-up. A total of 1876 implants in 406 patients was analysed. The survival rates of the implants varied from 100% to 72.4% (Table 4). The event rate for implant loss in the case of ≥ 6 implants and a splinted



**Table 3** Treatment procedures in the included studies.

| Study | Year of publication | Implants per patient | Pre-implant bone augmentation | Implant system | Anchorage design | Opposing dentition |
|--------------------------------------|---------------------|----------------------|--|---|------------------|---|
| Zou et al ⁴⁴ | 2013 | 4 | No | Straumann Standard SLA | Bar | # |
| | | 4 | No | Straumann Standard SLA | Locator | # |
| | | 4 | No | Straumann Standard SLA | Telescopic crown | # |
| Slot et al ⁴¹ | 2014 | 6 | No | Astra Tech AB | Bar | Natural teeth |
| | | 6 | Maxillary sinus floor augmentation | Straumann Standard SLA | Bar | Natural teeth |
| Slot et al ⁴³ | 2014 | 4 | Maxillary sinus floor augmentation | Straumann Standard SLA | Bar | Implant overdenture |
| | | 6 | Maxillary sinus floor augmentation | Straumann Standard SLA | Bar | Implant overdenture |
| Slot et al ⁴² | 2013 | 4 | No | Astra Tech AB | Bar | Implant overdenture |
| | | 6 | No | Astra Tech AB | Bar | Implant overdenture |
| El-Ghareeb et al ⁴⁰ | 2012 | 4 | Nasal floor augmentation | Brånemark MK III(20 implants) and Straumann Bone Level (4 implants) | Bar | All kinds of opposing dentition |
| Van Assche et al ³⁹ | 2012 | 6 | No | SLActive Standard Plus | Bar | All kinds of opposing dentition |
| Katsoulis et al ³⁸ | 2011 | 4 | No | Replace Select tapered | Bar | All kinds of opposing dentition |
| | | 5 | No | Replace Select tapered | Bar | All kinds of opposing dentition |
| | | 6 | No | Replace Select tapered | Bar | All kinds of opposing dentition |
| Mangano et al ³⁷ | 2011 | 4 | No | Leone implant system | Bar | # |
| Akça et al ³⁶ | 2010 | 4 | No | Straumann | Bar | All kinds of opposing dentition |
| Pieri et al ³⁵ | 2009 | 4 | No | PrimaConnex | Bar | All kinds of opposing dentition |
| | | 5 | No | PrimaConnex | Bar | All kinds of opposing dentition |
| Raghoobar et al ³⁴ | 2006 | 6–8 | Sinus floor augmentation and onlay block | Brånemark | Bar | # |
| Raghoobar et al ³³ | 2005 | 6 | Sinus floor augmentation | Brånemark | Bar | # |
| Payne et al ³² | 2004 | 3 | No | Brånemark | Ball | Two implant overdenture |
| | | 3 | No | Southern implant system | Ball | Two implant overdenture |
| Raghoobar et al ³¹ | 2003 | 6–8 | Sinus floor augmentation | Osseotite (3i) | Bar | All kinds of opposing dentition |
| Ferrigno et al ³⁰ | 2002 | 4–6 | Some | ITI | Bar | # |
| Zitzmann and Marinello ²⁸ | 2000 | 6–8 | # | # | Bar | # |
| Zitzmann and Marinello ²⁹ | 2000 | 6–8 | No graft procedures | Brånemark | Bar | # |
| Bergendal and Engquist ²⁷ | 1998 | 2–5 | No | Brånemark | Bar | All kinds of opposing dentition |
| | | 2–3 | No | Brånemark | Ball | All kinds of opposing dentition |
| Naert et al ²⁶ | 1998 | 4 | No | Brånemark | Bar | All kinds of opposing dentition |
| Watson et al ²² | 1997 | 3–4 | # | Brånemark | Bar | Natural teeth or implant supported prosthesis |
| Jemt et al ²¹ | 1996 | 3–4 | # | Brånemark | Bar | All kinds of opposing dentition |
| Hutton et al ²⁵ | 1995 | # | No | Brånemark | Bar | All kinds of opposing dentition |
| Jemt et al ²⁴ | 1994 | 4–6 | # | Brånemark | Bar | # |
| Johns et al ²³ | 1992 | # | No | Brånemark | Bar | All kinds of opposing dentition |

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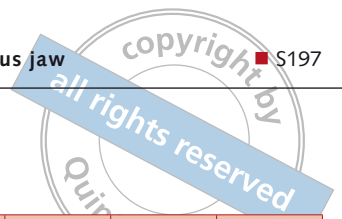


Table 4 Outcomes in the included studies.

| Study | Year of publication | No. of implants in study | No. of lost implants | No. of lost patients in study | Treatment (No. implants, mesostructure) | Survival rate implants (%) | Survival rate overdentures (%) | Change in marginal bone level (mean \pm SD; mm) | Gingival index (mean \pm SD) | Bleeding index (mean \pm SD) | Probing depth (mean \pm SD) |
|--------------------------------------|---------------------|--------------------------|----------------------|-------------------------------|---|----------------------------|--------------------------------|--|--------------------------------|--------------------------------|-------------------------------|
| Zou et al ⁴⁴ | 2013 | 40 | 0 | 0 | 4, bar | 100 | 100 | 1.0 (0.6) | 0.21 | 0.22 | 3.3 (0.7) |
| | | 40 | 0 | 0 | 4, locator | 100 | 100 | 0.9 (0.4) | 0.14 | 0.16 | 3.4 (0.5) |
| | | 40 | 0 | 0 | 4, telescopic crown | 100 | 100 | 0.9 (0.3) | 0.19 | 0.20 | 3.2 (0.8) |
| Slot et al ⁴¹ | 2014 | 150 | 3 | 0 | 6, bar | 98 | 100 | 0.22 | 0.2 | 0.3 | 4.3 |
| | | 150 | 1 | 0 | 6, bar | 99.3 | 100 | 0.5 | 0.1 | 0.6 | 4.3 |
| Slot et al ⁴³ | 2014 | 132 | 0 | 0 | 4, bar | 100 | 100 | 0.35 | 0 | 0 | 4.8 |
| | | 198 | 1 | 0 | 6, bar | 99.5 | 100 | 0.46 | 0 | 1 | 4.4 |
| Slot et al ⁴² | 2013 | 100 | 0 | 1 | 4, bar | 100 | 100 | 0.24 | 0.2 | 0.4 | 4.6 |
| | | 150 | 1 | 0 | 6, bar | 99.3 | 100 | 0.25 | 0.3 | 0.4 | 3.6 |
| El-Ghareeb et al ⁴⁰ | 2012 | 24 | 0 | 0 | 4, bar | 100 | 100 | # | # | # | # |
| Van Assche et al ³⁹ | 2012 | 72 | 1 | 0 | 6, bar | 98.6 | 100 | 1.3 | # | 0.28 | 3.4 |
| Katsoulis et al ³⁸ | 2011 | 88 | 1 | 0 | 4, bar | 98.9 | 100 | # | # | # | # |
| | | 25 | 0 | 0 | 5, bar | 100 | 100 | # | # | # | # |
| | | 6 | 0 | 0 | 6, bar | 100 | 100 | # | # | # | # |
| Mangano et al ³⁷ | 2011 | 152 | 4 | 0 | 4, bar | 97.4 | 100 | # | # | # | # |
| Akça et al ³⁶ | 2010 | 44 | 1 | # | 4, bar | 97.7 | 88 | 1.15 | 0.8 | 0.2 | # |
| Pieri et al ³⁵ | 2009 | 28 | 1 | 0 | 4, bar | 96.4 | 100 | # | # | # | # |
| | | 75 | 2 | 0 | 5, bar | 97.3 | 100 | # | # | # | # |
| Raghoobar et al ³⁴ | 2006 | 56 | 0 | 0 | 6–8, bar | 100 | 100 | # | # | # | # |
| Raghoobar et al ³³ | 2005 | 30 | 1 | 0 | 6, bar | 96.7 | 100 | # | # | # | # |
| Payne et al ³² | 2004 | 60 | 5 | 0 | 3, ball | 92 | # | # | # | # | # |
| | | 57 | 10 | 1 | 3, ball | 82 | # | # | # | # | # |
| Raghoobar et al ³¹ | 2003 | 68 | 3 | 0 | 6–8, bar | 95.6 | 100 | 0.3 (0.7) | 0.5 (0.7) | 0.7 (0.9) | 3.4 (1.3) |
| Ferrigno et al ³⁰ | 2002 | 114 | 3 | # | 6, bar | 92.2 (Milled bar) | 94.7 (Milled bar) | # | # | # | # |
| | | 64 | 6 | # | 4, bar | 86.9 (Dolder bar) | 87.5 (Dolder bar) | # | # | # | # |
| Zitzmann and Marinello ²⁸ | 2000 | # | # | 0 | 6–8, bar | # | # | # | # | # | # |
| Zitzmann and Marinello ²⁹ | 2000 | 71 | 4 | 0 | 6–8, bar | 94.4 | 100 | 0.92 | 54% (SD 26%) | # | # |
| Bergendal and Engquist ²⁷ | 1998 | 29 | 6 | # | 2–5, bar | 79 | 90 | 1.25 | # | # | # |
| | | 18 | 7 | # | 2–3, ball | 61 | 88 | 1.0 | # | # | # |
| Naert et al ²⁶ | 1998 | 53 | 6 | 6 | 4, bar | 88.6 | 85 | 0.5 | # | 0.2 (0.7) | 3.6 (0.9) |
| Watson et al ²² | 1997 | 117 | 30 | 14 | 3–4, bar | 72.4 | 77.9 | # | # | # | # |
| Jemt et al ²¹ | 1996 | 117 | 30 | 14 | 3–4, bar | 72.4 | 77.9 | 0.8 (0.8) | # | # | # |
| Hutton et al ²⁵ | 1995 | 117 | 29 | # | #, bar | 72.4 | 72.4 | # | # | # | # |
| Jemt et al ²⁴ | 1994 | 32 | 0 | 0 | 4–6, bar | 100 | 100 | Mesial side 0.30 (0.25) Distal side 0.34 (0.11) | # | # | # |
| Johns et al ²³ | 1992 | 117 | 21 | 5 | #, bar | 82.2 | 86.3 | 0.5 | # | # | # |

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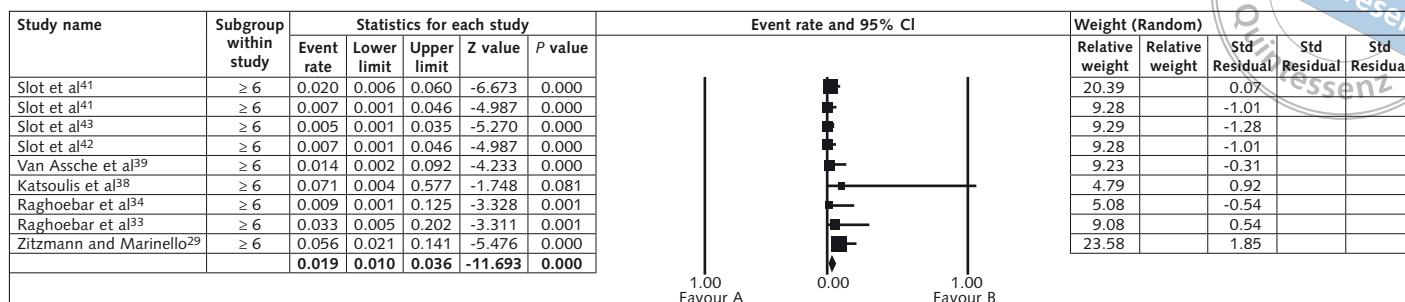
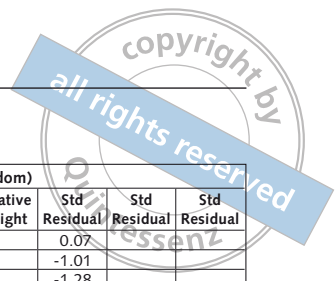


Fig 2 Meta-analysis of implant loss in case of ≥ 6 implants and a splinted superstructure. (When a study is mentioned twice, more than one implant system was analysed in that study. For details see Table 2.)

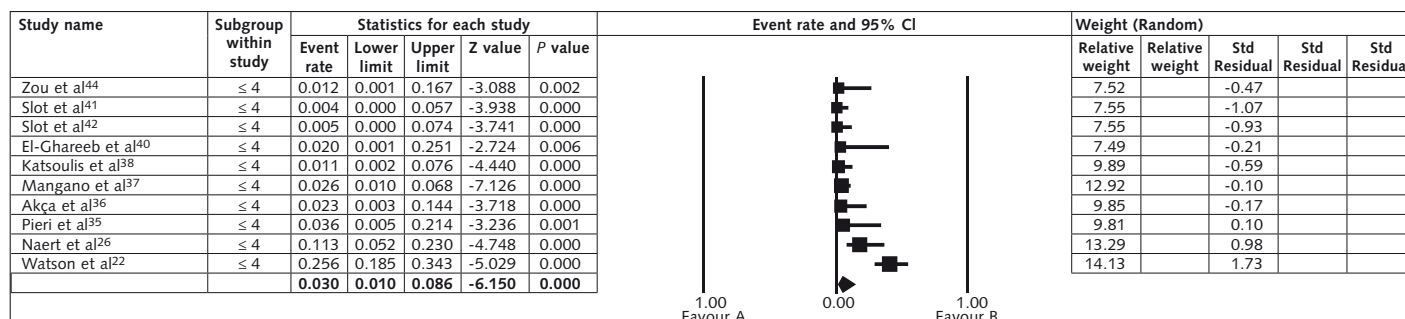


Fig 3 Meta-analysis of implant loss in case of ≤ 4 implants and a splinted superstructure.

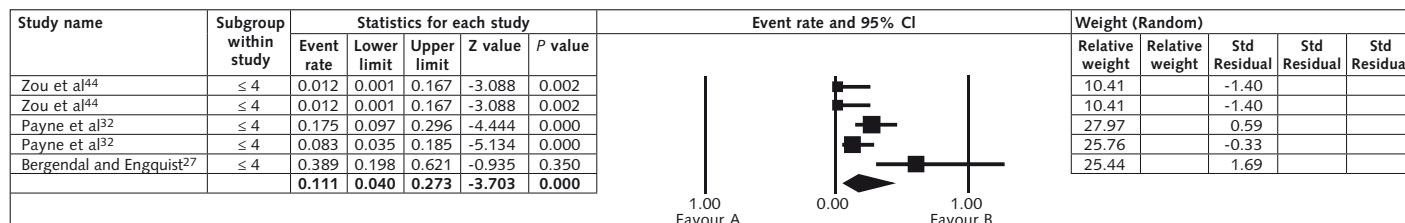


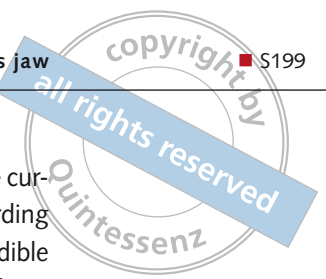
Fig 4 Meta-analysis of implant loss in case of ≤ 4 implants and a non-splinted superstructure. (When a study is mentioned twice more than one implant system was analysed in that study. For details see Table 2.)

anchorage was 0.019, which can be expressed as a survival rate of 98.1% per year (Fig 2). The event rate for implant loss in the case of ≤ 4 implants and a splinted anchorage was 0.030, which can be expressed as a survival rate of 97.0% per year (Fig 3). The event rate for implant loss in the case of ≤ 4 implants and a non-splinted anchorage was 0.111, which can be expressed as a survival rate of 88.9% per year (Fig 4).

■ Survival of maxillary overdentures

The survival of maxillary overdentures was defined as the percentage of overdentures initially placed that was still present at follow-up. Survival rates

of the overdentures varied from 100% to 77.9% (Table 4). The weighted meta-analysis (for person-years and for study size) for overdenture loss, expressed as event rates, in case of ≥ 6 implants and a splinted anchorage was 0.005 (95% CI [0.002 – 0.012]), which can be expressed as a survival rate of 99.5% per year. The event rate for overdenture loss in the case of ≤ 4 implants and a splinted anchorage was 0.031 (95% CI [0.013 – 0.076]), which can be expressed as a survival rate of 96.9% per year. The event rate for overdenture loss in the case of ≤ 4 implants and a non-splinted anchorage was 0.012 (95% CI [0.002 – 0.086]), which can be expressed as a survival rate of 98.8% per year²⁷.



■ Discussion

In contrast to the edentulous mandible, prospective studies with clinical and radiological baseline data reflecting the number of implants needed to support a maxillary overdenture, with an appropriate sampling frame, adequate sample size and sampling method are currently scarce. In addition, there is a shortage of RCTs to compare the outcome of specific questions related to the number of implants and design of the superstructure. In only two RCTs, the treatment outcome of 4 and 6 implants to support a maxillary denture was compared^{42,43}. In these RCTs no difference was noted between these treatment concepts after 1-year follow-up. All the other included publications provided data from convenience samples. Notwithstanding this drawback, on the basis of the available data we conclude that an implant-supported maxillary denture on at least 4 implants and provided with a bar anchorage is a proper treatment option for the edentulous maxilla, mainly because implant loss is considerably higher when the implant-denture is supported by < 4 implants.

By contrast and as mentioned before, there is a large body of evidence on which treatment concept is most suitable for the edentulous mandible. A 2-implant supported mandibular overdenture should be the minimum offered to edentulous patients as a first choice of treatment. The implant survival rate of mandibular overdentures is high, regardless of the number of implants¹⁵. Furthermore, there is evidence from systematic reviews and a large number of RCTs applying patient-based outcome assessments such as patients' satisfaction, oral-health related quality of life and in-depth qualitative interviews with patients that implant-supported mandibular overdentures have considerable benefits over conventional complete dentures¹⁴. It has to be mentioned, however, that the aforementioned recommended 2-implant supported mandibular overdenture treatment was based mainly on the results of studies that described implants placed in edentulous mandibles with a mandibular height in the symphysis region of at least 12 mm, and not in extremely atrophied jaws (mandibular height <12 mm). For the extremely resorbed mandible, there might be a need to modify this treatment concept. A treatment proposal for these very

atrophied mandibles based on the best evidence currently available in the literature is made². According to this proposal, in the extremely resorbed mandible (bone height and width ≥ 6 mm), 4 short implants could be placed if the soft tissues are in a good condition. Only in cases with a bone height of <6 mm, or when the soft tissue not in a good-enough condition to support an implant-supported mandibular denture, a bone augmentation procedure is advised.

In contrast to the excellent long-term implant and prosthodontic survival and success rates for implant-supported mandibular overdentures^{10,45,46}, several studies have described a higher number of implant failures and prosthodontic complications for implant-supported maxillary overdentures^{1,19,21,45}. Poor bone quality, low bone quantity, short implant length with reduced diameter and poor initial stability are problems observed in edentulous maxillae cases and may adhere to the higher risk of implant loss and loss of maxillary overdentures^{21,47}.

As reported, the 1-year implant survival rate in the case of ≥ 4 implants supplied with a bar anchorage is >95%, which is very promising and comparable to the concepts using 4 or 6 implants and a bar anchorage to support the maxillary denture⁴¹⁻⁴⁴. Reliable long-term data are not yet available. When losing an implant as part of 6-implant concept, a new surgical treatment procedure is usually not needed, as the overdenture can be adjusted. This is often not the case for the 4-implant approach, as with many of these patients a new implant has to be placed and a new suprastructure has to be made before the overdenture can be adjusted.

Progressive marginal bone loss is a predictor for future implant loss. Therefore, it is very important to analyse marginal bone loss in a standardised and reliable way. However, most studies used panoramic radiographs on which small changes in marginal bone loss are often not easy or not possible to assess. In the few studies that used standardised intraoral radiographs, marginal bone loss was less than 1.3 mm after 1 year, which is promising^{35,36,39,41-44}. Further studies are needed to truly rate the long-term marginal bone loss around maxillary implants.

Mucosa indices, bleeding indices and pocket probing depth provide insight into the health of the peri-implant soft tissues. In the studies covering this aspect, the soft tissues appeared relatively healthy,



although mucositis and gingival hyperplasia may occur around the implants and below the bar^{21,22}. Mucositis and gingival hyperplasia are usually reserved to conditions where the space between the bar and the oral mucosa or the space between the implants is limited. These conditions make proper oral hygiene difficult.

Future research concerning implant treatment of the edentulous maxilla should focus on long-term prospective clinical trials with detailed follow-up, in which clinical and radiographic aspects are analysed, restoration of function is assessed and patient satisfaction is scored. The current RCTs still only report on the 1-year follow-up data. Besides trials with overdentures, long-term RCTs comparing maxillary implant overdentures and fixed implant prostheses (e.g. costs, success rate, patient preference, and patient quality of life) are needed. Such comparisons are currently lacking. Only when all these factors are properly assessed will an evidence-based treatment concept for implant-supported maxillary dentures be found, thereby contributing to a higher level of care in this field.

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