DEALING WITH DENTAL IMPLANT FAILURES

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ABSTRACT

An implant-supported restoration offers a predictable treatment for tooth replacement. Reported success rates for dental implants are high. Nevertheless, failures that mandate immediate implant removal do occur. The consequences of implant removal jeopardize the clinician’s efforts to accomplish satisfactory function and esthetics. For the patient, this usually involves further cost and additional procedures. The aim of this paper is to describe different methods and treatment modalities to deal with dental implant failure. The main topics for discussion include identifying the failing implant, implants replacing failed implants at the exact site, and the use of other restorative options. When an implant fails, a tailor made treatment plan should be provided to each patient according to all relevant variables. Patients should be informed regarding all possible treatment modalities following implant failure and give their consent to the most appropriate treatment option for them.

Key words: Clinical studies. Endosseous dental implants. Implant restorations. Implant survival, success. Re-implantation. Fixed denture. Removable denture.

INTRODUCTION

An implant-supported restoration offers a predictable treatment for tooth replacement. Reported success rates for dental implants are high, however, there is still a paucity of data in the literature regarding follow-up of implants in function for at least 5 years or more. Nevertheless, failures that mandate immediate implant removal do occur. The consequences of implant removal jeopardize the clinician’s efforts to accomplish satisfactory function and esthetics. For the patient, this usually involves further cost and additional procedures.

Reported predictors for implant success and failure are generally divided into patient-related factors (e.g., general patient health status, smoking habits, quantity and quality of bone, oral hygiene maintenance, etc), implant characteristics (e.g., dimensions, coating, loading, etc), implant location, and clinician experience.

Cluster behavior can occur in implant failure. The finding that implant failures are not randomly distributed in the treated populations and that implant loss clusters in specific high-risk groups and individuals was examined in a literature review.

The aim of this paper is to describe different methods and treatment modalities to deal with dental implant failure.

Identifying failing implant

Success of dental implants is commonly defined by implant survival. Implant failure probably results from multifactorial process. There are various causes related to early (overheating, contamination and trauma during surgery, poor bone quantity and/or quality, lack of primary stability, and incorrect immediate load indication), and late (periimplantitis, occlusal trauma, and overloading) failure.

Ongoing marginal bone loss (MBL) could also put at risk implant survival in the long-term. In 1986, Albrektsson, et al. suggested success criteria for MBL, among other parameters. During the first year after abutment connection, 1 mm of MBL is allowed followed by 0.2 mm per year. Today, these criteria are still frequently referred to as the “gold standard” for implant success.

Recently, the abundance of data regarding MBL, and a better understanding of bone and soft tissue behavior around the implant neck and body, have shown these criteria to be inaccurate for today’s wide variety of implant systems.

An implant that causes clinical symptoms, such as continuous pain, mobility, etc, is considered faulty. However, MBL is rarely symptomatic but may endanger long-term implant survival. Although reports on the dynamics of MBL over time are incomplete, the MBL rate changes at different stages during the life of an implant. Given the accumulation of MBL data, calculations should not include a smooth polished neck portion. Long-term prognosis of an implant cannot be established based only on first year MBL calculations. Follow-up is essential to determine and predict
a future clinical course. Previously, we recommended that four clinically detectable MBL patterns be used for clinical follow-up and assessment. These hypothetical patterns of implant MBL after the first year were low rate MBL over the years (Albrektsson’s pattern), low rate MBL in the first few years followed by a rapid loss of bone support, high rate MBL in the first few years followed by almost no bone loss, and continuous high rate MBL leading to complete loss of bone support (For review see Schwartz-Arad, et al.).

Criteria for implant success should serve as an aid to clinical follow-up and to help evaluate the clinical outcomes of different implant systems in research. For clinical use, MBL assessment should be easy to apply using radiographs and should allow a quick gross comparison to previous data. Together with Albrektsson’s clinical parameters, it should help the clinician assess a given condition and predict its future clinical course, as well as help in decision making regarding additional tests/therapy (i.e., radiographs, occlusal analysis, prosthetic evaluation, surgical intervention, etc), frequency of follow-up, and hygiene appointments.

According to Albrektsson’s clinical parameters, in a low MBL rate during the first 3 years, the MBL pattern is still undetermined in an asymptomatic implant. Frequent follow-up is recommended to decide whether the implant is failing. Nevertheless, long-term follow-up is suggested for all MBL patterns.

It is essential to identify a failing implant in time to avoid continuous alveolar bone loss which might complicate the option of replacing the failed implant with a new one as well as impair the esthetic outcome of the area.

**Implant replacement**

The success of implants replacing failed ones at the exact site has been reported. Using the commercially pure titanium screw-shaped implants, it has been suggested that when an implant is lost, a flap should primarily cover the entrance to the site and after 9-12 months, a new implant can be replaced at that site. Evian and Cutler report immediately replacing 5 failed screw-type, commercially pure titanium implants with larger-diameter, hydroxyapatite-coated implants in the same sockets. They suggest that a 1-year healing period may not be necessary provided the socket can be prepared to eliminate thread grooves and invasive soft tissue; the implant replacement is larger in diameter than the original implant; and sufficient available bone remains for the procedures. Recently, the implant failure rate was compared between a machined surface and a TiUnite surface used to replace failing implants. Of the 29 machined-surface implants replaced by implants with the same surface, 6 failed (79.4% survival rate) compared to the 19 machined-surface implants replaced by TiUnite surface implants where only 1 failed. Of the 10 TiUnite-surface implants replaced by implants with the same surface, none failed. The difference in failure rate between machined-surface and TiUnite replacement implants was statistically significant.

In a study that assessed survival and success rates of single dental implants replacing a previously failed implant at the same location, an overall survival rate of 71% was reported with a mean follow-up of 19.4±11.4 months.

Replacement of a failing implant involves the challenge of achieving osseointegration in a compromised bone site. When treatment cost and additional procedures to the patient are considered, the clinician needs information regarding the predictability of replacing a failed implant. This information should be discussed with the patient for informed consent for the subsequent attempt.

There is still a lack of sufficient evidence-based data regarding failed implant replacement. Metucci removal of granulation tissue on the failed implant site and the use of wider implants with improved surfaces could improve the outcome of re-implantation. Further research with a large cohort for a long follow-up period is warranted.

An implant that replaces a previously failed one could serve as a predictable procedure with reasonable survival rates. However, these survival rates are lower than the rates reported for first attempt single implant placement. Clinicians should remember that once an implant has failed, replacement of that implant is subjected to at least all the initial factors that led to the failure.

**Other restorative options**

**Short arch**

When planning implant rehabilitation or when facing implant failure, one should always refer to the question: How many teeth are necessary for adequate function or what dentition assures oral function? In some instances, the treated area can remain edentulous and this should be considered as an option. A key indicator of oral health status is the number of teeth. In 1992, the World Health Organization (WHO) stated that throughout life, the retention of a functional, esthetic, natural dentition of 20 teeth, without requiring prostheses, should be the treatment goal for oral health. Therefore, 20 teeth have been used as an operative expression for a functional natural dentition in epidemiological studies.

After extensive review of the literature, Elias and Shelmiah concluded that to satisfy oral functional needs a complete dentition is not necessary, which is in accordance with others who suggest that middle-aged and older adults have sufficient oral function with 20 natural teeth, and question the need to replace missing molars. However, the demand for tooth replacement was assessed under normative and theoretical conditions, rather than among patients who had experienced tooth loss.

The relationship between dentition and oral function has been evaluated in a review, and concludes that the World Health Organization goal for the year 2000, namely to maintain a natural dentition of not less than 20 teeth throughout life, is substantiated by the current literature since this proposed dentition will assure an acceptable level of oral function. This should also be remembered when dealing with implant failure.

**Fixed partial denture**

The alternative use of fixed partial denture (FPD), if applicable, is another treatment modality. Recently, a
thorough systematic review was conducted that analyzed and compared the survival and success rates of different designs of tooth and implant-supported fixed reconstructions and assessed the incidence of biological and technical complications of FPDs and dental implants. The incidence of technical complications was significantly higher for the implant-supported reconstructions compared with tooth-supported FPDs. In another review that analyzed tooth loss and evaluated the longevity of healthy teeth and teeth compromised by diseases and influenced by therapy and oral implants, found that unless affected by oral diseases or service interventions, teeth will last for life. Numerous retained teeth could be an indicator of positive oral health behavior throughout life. Tooth longevity is largely dependent on the health status of the periodontium, pulp or periapical region, and extent of reconstructions. Multiple risks lead to a critical appraisal of the value of a tooth.

Oral implants when evaluated after 10 years of service present with a longevity that does not surpass that of even compromised, but successfully treated and maintained teeth.

Removable denture

Removable partial dentures (RPDs) are still extensively used for the restoration of partially edentulous patients. However, these prostheses have been associated with poor patient acceptance, compromised function and esthetics, and increased risk for caries and periodontal disease. Some even consider the art of removable prosthodontics as obsolete. However, there is an increased need for the management of partially edentulous patients, which is due to the increase in life expectancy and the well-documented decline in tooth loss and total edentulism in the US over the past several decades. Adults in the US retain approximately 2.0 more teeth every decade, which explains the unmet need of 516 million chairside hours estimated for prosthetic treatment alone in the US by 2010, and the increase to 5060 million hours by 2020. Approximately 66% of this unmet need is for FPDs and 34% for RPDs.

Despite the obvious need for RPDs, a detailed search of the dental literature failed to elicit strong evidence-based indications for treating the partially edentulous patient with a conventional clasp-retained RPD. When implants fail or are not an option and economic considerations preclude extensive fixed restorations, RPDs are a valid treatment alternative.

The combination of dental implants to support the RPD may alleviate some of the problems associated with the conventional RPDs.

Implant tooth-supported removable partial denture (RPD)

The problematic long-term clinical experience of restoring partially edentulous patients with RPDs in the era when implants are predictably used for the same patient group suggests the use of implants in combination with RPDs. Implants are used to improve the RPD support, enhance retention and stability, preserve the residual ridge underneath the denture base, reduce the stress applied on the abutment teeth, eliminate the need for un-esthetic clasp assemblies, and modify unfavorable arch configurations. Generally, RPDs are still needed in cases of un-replaced failed implants, or where economic, systemic, or local anatomic conditions preclude the use of extensive rehabilitation with fixed implant-supported restorations.

Laboratory and clinical studies show the effectiveness of implant-supported RPDs. In model laboratory studies, distally-placed implants supporting a mandibular Kennedy Class I RPDs prevented displacement of the distal extension implant-supported RPDs, improved occlusal support, and decreased the pressure on soft tissues compared to conventional RPDs.

In a retrospective clinical study, 10 patients were treated with uni- and bi-lateral mandibular distal-extension RPDs supported by 16 posterior implants. Implants were used as either vertical stops to enhance the prosthesis support or with resilient retentive elements. There was consistently increased satisfaction in all patients, minimal component wear, no radiographic evidence of excessive bone loss around the implants, and stable peri-implant soft tissues. In another study, 15 partially edentulous patients with an unfavorable number and distribution of abutments were treated with implant-supported RPDs. The partially edentulous arch configuration was modified by placing 33 implants into strategic sites. An implant survival rate of 100% was reported, with only minor prosthetic complications and significantly improved patient satisfaction.

In a case series, 23 implant-supported RPD were placed in 44 implants during a 10-year period (1996-2005). Maxillary restorations were provided to 13 patients and mandibular to 10 patients. Before implant placement, the most prevalent arch configuration was Kennedy Class I in the maxilla (6 patients) followed by Kennedy Class II in the mandible (4 patients). Arch configuration was modified by implant placement in 6 (26.1%) patients. Survival rate was 95.5%; only two implants did not survive, both in a heavy smoker with pre-existing periodontal disease. During follow-up, only one abutment tooth was lost. All other abutments remained in function with no need for re-treatment during the last recall. All patients were highly satisfied with the restoration. According to our findings, we suggest that implant-supported RPD could serve as a long-term predictable treatment modality. Nevertheless, a long-term multi-center study is recommended to evaluate the success of this treatment modality in a larger patient sample. Prospective clinical studies should focus on abutment longevity and need for prosthesis maintenance.

CONCLUSIONS

Implant therapy has become common practice and will probably gain in popularity during the next several years. This implies that dental professionals will have to deal more
with implant failure and related complications. When an implant fails, a tailor made treatment plan should be provided to each patient according to all relevant variables. Patients should be informed regarding all possible treatment modalities after implant failure and give their consent to the most appropriate treatment option for them.

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