

PRELIMINARY STUDY ON THE SUCCESS-FAILURE BALANCE IN IMPLANT THERAPY

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Abstract:

Implant therapy has been shown to be predictable, effective and reliable for the replacement of missing teeth in edentulous and partially edentulous patients. Prosthodontic rehabilitation with dental implants requires accurate implant placement for predictable functional and aesthetic outcomes. Implant dentistry has developed numerous advancements in technology, materials, techniques and concepts to achieve the desired beneficial clinical results. Clinical evidence has demonstrated long-term (> 10 years) survival rates up to 95%. Despite the high success and survival rates of dental implants, failures can and do occur. There are certain factors such as smoking, history of periodontal disease which have been associated with higher rates of implant failure. Furthermore, implant site characteristics such as decreased quantity and/or quality of bone, infected sites and anatomic localisation such as the posterior maxilla have also been correlated to decreased implant survival.

Key words: dental implants, hard tissue, infection, interdental space, local risk factors, soft tissue.

INTRODUCTION

The oral cavity has been described as a mirror that reflects the health of the body [1]. A correct diagnosis, elimination of the causes and reduction of modifiable risk factors are paramount for successful prevention and treatment of oral diseases [2]. Nowadays, the pathology of the oral

cavity, especially edentation, represents a major public health problem [3].

The oral cavity is colonized by a complex microbiota that grows and lives as diverse biofilms on all mucosal and dental surfaces. Oral biofilms on dental hard and soft tissues are the main cause of dental

diseases, including caries and periodontal disease [4]. The low pH medium is a primary factor of decays that occurs through saliva acidification as result of alimentary and hygiene individual behaviors like food diet and tooth brushing. The modifications of salivary pH could be also caused by the influence of bacteria, enzymes, hormones and other factors [5].

Concerning the etiology, location, extent or topography, edentation is considered a pathological entity that, through evolution and complications, causes serious imbalances in the stomatognathic system [6]. Teeth are extracted as a consequence of an oral complication, following the interaction between the patient and the dentist, the dentist's ability to provide treatment to sustain the functionality of the teeth and depending on the patient's preference [7]. Actually, the dental medicine is governed by special aesthetic demands based on avant-garde techniques and technologies in conjunction with the type of biomaterials used. [8].

The use of dental implants is considered a revolution in modern dentistry [9,10]. Osseointegrated dental implants have represented a breakthrough in clinical practice for replacing missing teeth and supporting prosthetic reconstructions in edentulous areas [11].

The use of dental implants to aid in the support of restorations replacing missing teeth has been reported in the literature dating back to the early 1960 [12,13]. Historically, dental implant treatments have had mixed results with regard to survival of the implants and prostheses. In the context of society development, the past two decades have seen theoretical and practical interest of manufacturers, researchers and

clinicians to improve the success of implant treatment outcomes through evolution in implant design, materials and clinical procedure [13, 14].

While the patients is usually concerned with esthetics and function, dental professionals expect success regarding biological and mechanical stability and the facilitation of oral hygiene [10,15]. There is consensus among authors that the success of dental implant treatment depends on the presence and maintenance of surrounding bone mainly in the bone crests area. However, one of the major challenges encountered in implantology is the process of bone resorption around the implant after insertion or during its use. In the literature, bone resorption of approximately 1.2 mm in height during the first year of function is reported with 0.1 mm more resorption for every subsequent year [10,17,18 16,17]. This loss with a V or U shape has been called saucerization [10, 18].

In 1986, Albrektsson et al establish the following criteria for implant success [10,19]. The implant should have no mobility and demonstrate no radiolucent areas radiographically, annual vertical bone loss after the first year should be less than 0.2 mm and there should be no persistent and or irreversible symptoms. The most common parameter used in clinical reports is the survival rate, indicating whether the dental implant is physically in the mouth or has been removed [10,20]. However with this method, implants that should be removed owing to pain or illness may be retained and erroneously considered successful.

In 1993, an implant quality of health scale was created by James and developed by Misch [10, 22, 22]. This scale was later modified at the International Congress of

Oral Implantologist's Pisa Consensus Conference in 2007, presenting four clinical categories that contain condition of success survival and failure of the implant. Survival can be divided into two categories: satisfactory survival which describes implants with less than ideal condition, but for which there is no need for clinical intervention and compromised survival includes implants with less than ideal conditions required clinical treatment to reduce the risk of implants that require removal or that have been lost. Implant success is a term used to describe clinical conditions and must include at least 12-month period for implants serving as prosthetic abutments. Early success is suggested for implants that are retained for a period of one to three years, intermediate success for three to seven years and long-term success for a period longer than seven years. In this new approach, pain, mobility, radiographic bone loss, probing depth and periimplant disease are evaluated [10,23].

Regarding periimplant disease, since the bone loss caused by stress or bacteria leads to the deepening of the sulcular gap and decreases oxygen tension, anaerobic bacteria become the primary promoter of continuous bone loss. Exudate or an abscess around an implant indicates exacerbation of periimplant disease and possibly accelerated bone loss. Exudate persisting for more than one or two weeks normally requires surgical intervention in the periimplant area to eliminate the etiological factors. The reduced bone height after the exudate episode exposes the implant to secondary occlusal trauma. The surgeon specialist should re-evaluate and reduce the stress factors for the new bone condition to improve the performance in the long term [10,23].

The aim of our 15-year study is to determine the effects of factors with local potential risk on implant survival and success (primary outcomes) as well as on mucosal recession, bleeding on probing and proximal marginal bone loss (secondary outcomes). The following local risk factors were considered: interdental space, infected sites, soft tissue thickness, width of keratinized soft tissue, bone density and implant stability. As with any study involving the oral cavity, oral hygiene is a key variable. There is substantial evidence that poor hygiene is associated with periimplant [24,25].

The following clinical and radiographic criteria proposed by Albrektsson et al and adapted by Buser et al and Karoussis et al were used to define implant success [26,27,28]: absence of mobility, absence of persistent subjective complaints (pain, foreign-body sensation and/or dysesthesia, absence of recurrent peri-implant infection with suppuration, absence of a continuous radiolucency around the implant, no pocket probing depth >5 mm, no bleeding on probing, annual vertical bone loss after the first year of service not exceeding 0,2 mm (mesially or distal) [26,27,29,30,31]. Several secondary outcomes were also considered. These included mucosal recession, bleeding on probing and proximal marginal bone loss. Although these outcomes individually were not directly related to implant survival or success, they were determined to be significant when considering outcomes in the esthetic zone.

It is important to differentiate between the meaning of Survival and Success. „Implant survival,, means that implants are still in the mouth at the time

of examination, regardless of the state of the prosthesis or patient satisfaction. A nonfunctional implant requiring additional treatment is counted in the surviving group while „Implant success,, means that implants are not only in the mouth but are also functional and satisfactory.

MATERIAL AND METHODS

This study included 624 patients who received 1127 implants and prostheses on implants in a private clinic in the period 2006-2021. These included completely and partially edentulous arches as well as single-tooth replacements. All the prosthetic bridges performed were fixed. Of the 624 patients studied, 345 (55.28%) are women and 279 (44,71%) are men.

After a complete medical and dental history, through clinical and radiographic evaluations including as indicated, periapical, panoramic, conventional and computerized tomographic films, diagnostic casts and consultation with the restorative dentist were carried out. Patients were presented with treatment options and advised of the potential problems associated with dental implants and an informed consent form was signed. Patients were not accepted for implant surgery if they were medically contraindicated for elective surgery or had unrealistic expectations.

All implants were placed under sterile conditions following this protocol: the patient was administrated intravenous sedation using fentanyl citrate 0,5 mg/ml injection, diazepam 5 mg/ml injection, as well nitrous oxide, oxygen inhalation. This was in addition to articaine hydrochloride 4% and epinephrine bitartrate 1:100 000 local anesthesia that was administrated in

both block and infiltration technique. They also received antibiotics Augmentin 24 hours prior to surgery and 6 days thereafter (1mg twice a day), corticosteroid medication (Medrol) for control of the inflammatory response was given daily in a regressive mode (15 mg at surgery, 10 mg on the first 2 days postoperatively and 5 mg in days 3 and 4 postoperatively), antiinflammatory medication Arcoxia (Merk) 120 mg once day, was administrated on day 5 postoperatively every 12 hours. Anagesics were given on the day of surgery and postoperatively for the first 3 days as needed. Because the bone density allowed us, the insertion torque of the implants was >50 N.cm. Sinus lifts were performed in patients who did not have enough space for implant insertion, along with bone additions with Bio-Oss (Geistlich Pharma AG) and resorbable membrane Hypro-Sorb F.

All patient records, histories and radiographs were reviewed; all implant locations were recorded appropriately as posterior maxilla, anterior maxilla, posterior mandible or anterior mandible and all implant failure were recorded.

The succes and survival rates of the implants were analyzed based on the criteria of the Pisa Consensus Conference according to the following clinical parameters: pain (absent, absent in function, sensitivity in function, pain in function), mobility (present or absent), probing depth (PD), bleeding on probing (BOP), exudate (absent, with exudate history, with uncontrolled exudate) and radiographic bone loss. The PD and BOP measurements were taken at four aspects of each implant: mesial (M), distal (D), buccal (B) and lingual/palatal (L/P).

For the assesment of radiographic bone loss, a periapical radiograph using the

bisecting angle technique was performed at the time of patient recall. According to the criteria, the implants were classified as successful or failed.

Biological and prosthetic complications such as periimplant mucositis, periimplantitis or any mechanical and prosthetic complication were also evaluated. Patient with BOP or positive suppuration, a PD greater than 5 mm and radiographic bone were diagnosed as having periimplantitis.

Table 1. Number of patients with dental implant

Patients with implants					
women		men		total	
number	%	number	%	number	%
345	55.28	279	44.71	624	100

Table 2. Number of implants by placement area

Number of implants and placement area									
Anterior and posterior maxilla				Anterior and posterior mandible				Total	
women		men		women		men			
number	%	number	%	number	%	number	%	number	%
234	20.76	170	15.08	381	33.81	342	20.35	1127	100

The implants used in this study were: ANKYLOS DENT SPLY (32 implants-3,01%), ALPHA BIO (173-16,27%), MIS BIOCROM (63-5,92%), AB (TITAN IMPLANT) (341-32,07%), BREDENT BLUE SKY (536-50,42%), ADIN (1-0,094%), TUFF (1-0,094%), NORIS (2-0,188%).

Of the 624 patients, only 52 patients needed sinus lift interventions, of which only 29 required bone additions. The materials used were: BIO-OSS (Geistlich Pharma AG), NANO-BONE (Artoss Gmgh), HYPRO-OSS (natural

RESULTS

During the study period between 2006-2021 was analyzed a number of 624 patients who met the inclusion criteria for this study. At these patients we encountered a number of 1127 implants. The age range of these patients was between 20-75 years.

hydroxyapatite and atelocalgen composite for bone substitution), KASIOS (TCP DENTAL HP) (calcium phosphate ceramic bone filling) and collagen resorbable membrane: HYPRO-SORB F (bilayer bioabsorbable barrier for guided tissue and bone regeneration), R.T.R cone (SEPTODONT) (resorbable tissue replacement).

The 29 bone additions were performed in the following cases: one case of bone addition and resorbable membrane immediately, two cases of bone additions and immediate placement of the implant, two

cases of bone addition+extraction and implant placement, two cases of extractions with curettage and bone addition and immediate placement of the implant, one case of periapical resection and immediate placement of the implant, 21 cases of extractions with bone additions and placement of implants immediately post-extraction.

Of the 52 sinus lift procedures: 26 cases were simple sinus lift, one case of bilateral sinus lift, one case of sinus lift+cystectomy+implant, 10 cases of sinus lift+bone additions+resorbable membranes+immediate placement implants, two cases of sinus lift+extractions+implant placed immediately post-extraction, five cases of sinus lift+implant placed immediately+membrane on implant, five cases of sinus lift+immediate implant placement, two cases of sinus lift+bone addition+membrane resorbable+extraction.

The size of the implants used in the jaw were: 3,3/10 mm, 3,5/10 mm, 3,5/11 mm, 3,75/8 mm, 3,75/13 mm, 3,75/16 mm, 3,75/10 mm, 4,8/8 mm and in the mandible were: 3,5/14 mm, 3,5/8 mm, 3,75/13 mm, 3,5/10 mm, 3,75/11,5 mm, 3,75/10 mm.

Based on the criteria for failure outlined above, 28 implants were rejected, of which 7 in the jaw and 21 in the mandible.

Of the 28 cases of rejection: 5 cases were rejected at 7 days with pain, mobility positive, positive depth probing (PD) as well as positive probing bleeding (BOP). These were suppressed and bone addition was affected with Bio-Oss. Another 3 cases of implants were rejected after 10 days showing pain, mobility, positive probing

depth (PD), positive bleeding on probing (BOP), exudate and the presence of bone loss on radiography. They were suppressed and bone addition was affected with Bio-Oss. Another 6 cases suffered rejection at 30 days with mobility, positive probing depth (PD), positive bleeding on probing (BOP), radiographic bone loss present. They were also suppressed and bone addition was performed with Bio-Oss. Another 4 cases were rejected at 21 days for the same reasons, two cases were rejected at 10 days being present in all these cases: mobility, pain, positive probing depth (PD), bleeding on probing (BOP) as well as bone loss radiograph. For cases were rejected immediately due to suppurative phenomena, pain, mobility, positive probing depth (PD), bleeding on positive probing (BOP) exposing the vestibular wall. Two cases suffered rejection 32 days after implantation due to mobility, positive depth probing (PD), positive bleeding on probing (BOP), exudate presence and radiographic bone loss. In these two cases, gingival curettage and addition of Bio-Oss were performed. Another case was that of an implant rejected after 14 days due to the presence of mobility, inflammatory exudate, positive PD and positive BOP with vestibular wall exposure. The last rejection appeared 90 days after implantation due to mobility, PD positive, BOP positive, exudate presence and radiographic bone loss. It was suppressed and Bio-Oss was added.

DISCUSSION

This study aimed to evaluate the success, survival and failure rates of implants based on the implant quality of health scale developed at the Pisa

Consensus Conference. The success category describes optimal conditions, the failure category includes implants that should or could be removed.

In this study, patients who underwent implant rejection first observed mobility, PD positive and BOP positive, then the presence of inflammatory exudate and then pain. So among the first signs that alerted patients were: mobility, positive PD, positive BOP, inflammatory exudate and then was pain and final confirmation was made based on radiographic bone loss.

The literature shows a lower success rate of implants placed in the maxilla compared with the mandible, a fact that is related to the lower density of the maxillary bone [32,33]. The residual bone height becomes insufficient owing to the loss of alveolar bone. However, the molar area in both the maxilla and mandible displays substantial bone deficiency owing to increased occlusal forces, increasing the failure rate of implants in this area [32]. In type IV bone, the cortical bone is very thin and the lack of dense bone makes it difficult to achieve adequate stability. The mandibular retromolar area and the maxillary molar region are formed by low-quality bone, while implants placed in the anterior mandible area have high success

rates owing to increased cortical bone. In this study, there was no difference between implants placed in different regions of the jaws regarding their success or survival corroborating the findings of Kim et al [34,35].

In our study, various implant systems were analyzed without differences in the success rate or acceptable survival rate, corroborating the findings of Ferrigno et al and Telleman et al who found similar results for the survival of different types of implant designs [36,37,38]. In a literature review, Opperman et al concluded that regarding implant survival, there are no types surfaces or implant systems that present clear advantages over others [39,40].

In summary, in the present study, of the 1127 implants made, only 28 (2.48%) were rejections.

CONCLUSION

Within the limitation of this study, the data suggested that the implant success rate does not seem to be related to factors like age, sex, systemic disease, macroscopic characteristics or area in which the implant was placed. This study can be considered preliminary and provides the basis for the design of further studies.

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