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POLYAMIDE VS SILK SUTURES IN THE HEALING OF POSTEXTRACTION SOCKETS: A SPLIT MOUTH STUDY

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SUMMARY

Objective. The aim of the present work is to compare polyamide sutures with traditional silk sutures after tooth extraction.

Methods. The investigators implemented a split-mouth study, in 50 healthy patients undergoing dental extractions. Follow-up included one post-extraction checkup at 7 days for the suture removal. The outcome variables were the Healing Index, the socket area, postsurgical complications and outcome of a patient questionnaire. Pairs of correlated continuous variables were analyzed with the Wilcoxon test, independent continuous variables with the Mann-Whitney test, and categorical variables with the $\chi 2$ test or Fisher test.

Results. From September 2015 to September 2016, 50 patients underwent contemporary bilateral extractions of homologous teeth. The treatment-*versus*-control postoperative comparison showed that polyamide sutures resulted in significantly smaller residual socket areas and better Healing Indices at day 7. The patients' questionnaire outcomes were unanimously in favor of polyamide sutures.

Conclusions. The primary goals of this study were to measure socket reduction and the Healing Index (HI) at a 7 day postextraction checkup and to assess patients' satisfaction based on answers to a questionnaire. Based on the results here reported, the polyamide sutures performed better than the silk sutures.

Key words: silk sutures, polyamide sutures, sutures, postextraction sockets.



Introduction

Depending on the structure, sutures are classified as monofil or polyfil, while they are absorbable or nonabsorbable owing to their chemical composition. Among the absorbable materials polydioxanone, polyglicolic acid and poly-Llactide, poly-e-caprolactone and polyglactin are used. Natural products as silk (polyfil) belong to the group of non-absorbable materials. Synthetic non-absorbable suture products consist, for instance, of polyamides, polypropylene and poly-

ester. Polyamides are available as monofil, polyfil and pseudopolyfil (polyamide fibers endowed with a polyamidic coat) threads.

Suture materials are prone to the adhesion of bacteria, which represent potential risk factors for wound healing. Indeed, the adhered bacteria, especially the anaerobic ones, can act as a focus of odontogenic infections. Fusobacteria, peptostreptococci, prevotella, porphyromonas, *Streptococcus intermedius, S. anginosus*, and bacteroides species are usually identified in odontogenic infections (1). Recent studies have shown that bacteremia can result from the re-

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moval of sutures (2, 3). Among the others, Otten et al. (4) investigated the bacterial colonization of a resorbable copolymer of glycolide and epsilon-caprolactone and non-resorbable polyamide monofilament sutures used in intraoral dentoalveolar surgery. The *in vivo* findings suggested that, in comparison with the resorbable sutures, more pathogens were present on non-resorbable sutures. Based on the *in vitro* study the colonization rate of *Streptococcus intermedius* on both sutures was similar, whilst *Prevotella intermedia* grew much better on polyamid than on monocryl.

Mechanical properties of these materials vary by the composition. The tensile and knot security properties of several resorbable and non-resorbable sutures was studied by Kim et al. (5) leading to the knowledge that no single suture material possesses all of the requirements depending on the type and caliber.

Sutures are applied in many medical areas, but, in dentistry, a few papers deal with these materials. For instance, polyamide suture was compared with polyglactin suture (6) in oral implant surgery. No difference could be found between the two suture groups, although a higher incidence of complications was found in the absorbable suture group. Polyamide sutures have been successfully used for periodontal surgery (7, 8).

To the Authors' knowledge, this study is the first comparing silk and polyamide sutures and focusing on the quality of the wound healing in a large number of patients who received dental extractions.

Materials and methods

Research design and management

To address the research purpose, a split-mouth study was designed and implemented. Post-extractive sockets were sutured either with polyamide sutures (Polimid, Swden & Martina Due Carrare (PD), Italy; investigation site), or with silk suture (Silkam, Silkam Non Absorbable Aesculap Aesculap, Inc. Center Valley, PA; control site). The choice of the socket was randomized recurring to a software.

The study population consisted entirely of patients referred to the Department of Surgery at the Dental School of the University of Turin from September 2012 to September 2013. The inclusion criteria were: good general health conditions, attendance of the preoperative professional hygiene program; need for bilateral extractions of homologous teeth; compliance to the study protocol. The exclusion criteria were platelet dysfunction, thrombocytopenia or treatment with corticosteroids.

Fifty patients satisfying all the inclusion and exclusion criteria were enrolled in the study. The study was planned and performed in accordance with the Declaration of Helsinki and was approved by the Ethics Committee of Dental School (Ref. Numb. CIR141113).

Surgical protocol

At enrolment, the patients were registered in a computerized clinical file and attended a program of professional dental hygiene. After a thorough clinical and radiologic evaluation, a digital case history for each patient was filed containing information on age, gender, smoking habits, alcohol consumption, systemic pathologies, the use of any drugs, oral and dental parameters such as periodontal screening and recording code. The possible presence of infections, carious processes, or periodontal affection were also noted as the difficulty of extraction (based on crown integrity, root morphology, space, and patient compliance).

Depending on the dental arch, alveolar nerve block infiltration was administered with either local or regional anesthesia, using 2% mepivacaine, which does not cause restriction of the blood supply, owing to the absence of epinephrine. To prevent interference with the healing process, no intraligamentous or intrapapillary infiltrations were performed. Non-traumatic bilateral extractions were accomplished without ele-

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vation of full-thickness flaps to preserve the bone ridges and soft tissue at the 2 sites. Surgery was performed on the treatment and control sides at the same time.

After removed granulation tissue, a millimeter probe was used to measure the socket diameters (maximum mesiodistal [MD] and buccolingual extensions [BL]). Accurate postoperative recommendations were provided to the patients. No pharmacologic therapy was prescribed, and recourse to antibiotics was deemed a negative outcome as a sign of postoperative complications from infection.

After the extractions, each patient was recalled at day 7. Measurements of the 2 variables – MD, BL – was conducted to determine the socket area (SA). The residual socket area (RSA) i.e., the fraction of the postextraction original cavity still open at the follow-up session (t = 7), was computed as the ratio of (MD \times BL)_t to the value (MD \times BL)_{t=0}, measured after extraction (perfect closure corresponds to RSV = 0.00).

The maturation and quality of regenerating tissues were evaluated using a modified version of the HI of Masse et al. (9-11), originally developed to evaluate healing with primary closure after periodontal surgery as previously reported by Mozzati et al. in 2013 (12). The modified HI, adapted to estimate socket healing without primary closure, involved 3 scoring levels for each of the 4 parameters considered: tissue color (1 = 100% of gingiva pink; 2 = <50% of gingiva red, hyperemic, movable; 3 = 50% of gingiva red, hyperemic, movable), color and consistency of the healing tissue (1 = close grained, pink; 2 =soft, red; 3 = fragile, greenish or grayish), suppuration (1 = absent; 2 = absent but pronouncedamount of plaque around socket walls; 3 = pronounced), and bleeding (1 = absent; 2 = induced)by palpation; 3 = spontaneous). Thus, the scoring scale ranged from 4, corresponding to excellent healing, to 12, indicating severely impaired healing.

Pain was evaluated by the patients using a VAS, which distinguished the study and control sockets, for 7 consecutive days, with day 0 considered the day of extraction, after resolution of the effects of the anesthesia.

The patients' general satisfaction was evaluated with a questionnaire administered after extraction. Patients were asked to state any differences noticed in bleeding or pain between the 2 sites and to state which technique they would choose for a future extraction.

Study variables

The primary predictor variable in the trial was treatment group status (study or control socket). The primary outcome variables were socket area, HI, and postsurgical complications (mainly the need for late follow-up and reoperations). The secondary outcome variable was the patient satisfaction. A third category of variables described the anamnesis of the sample (age, gender, smoking habits) and the dental characteristics (tooth vitality, caries, periodontal diseases, granuloma, and pain).

Statistical methods

The results included continuous and categorical variables. The former are reported as mean and standard deviation. Non parametric tests were used: the Wilcoxon signed-rank test for comparisons of 2 correlated samples involving matched pairs and the Mann-Whitney test for comparisons of 2 independent distributions.

Categorical variables, reported as count and percentage, were arranged in cross-correlation tables and compared using the χ^2 test with the Yates correction when all expected values were higher than 5 or the Fisher test. Statistical significance corresponded to a probability less than .05 that differences could be ascribed to chance.

Results

The patients enrolled in the study were 50 (36% female) with a mean age of 52 years (SD= 14). The complete absence of any complications

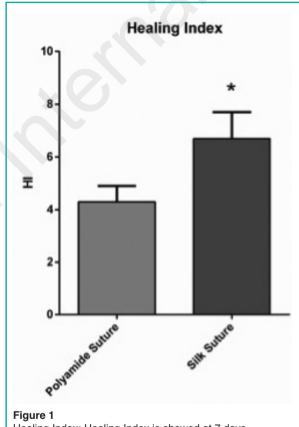


Days from extraction			
	Socket Areas		RSA
	Day 0	Day 7	Day 7
Polyamide suture	32.5±27.8	5.2±12.3	0.09±0.1
Silk suture	28.9±20.6	10.1±15.1	0.29±0.3
t-test value	0.234	0.041	328* 10-6

from surgery, such as infection, excessive bleeding, or alveolar osteitis, was a noteworthy result, which held true for both the study and control sides. Data concerning the healing of the sockets as for their progressive decreases in areas are presented in Table 1, while the values of HI are showed in Figure 1. Comparisons between values relative to the study and control sides showed better healing and faster socket closure for the wounds sutured with polyamide at day 7. Figure 2 shows the HI distribution. The patients' questionnaire outcomes were unanimously in favor of polyamide sutures.

Discussion

In the mouth more than in other districts, sutures should avoid or limit bacterial adhesion and proliferation. The present study was designed to clinically test the validity of polyamide sutures against silk ones: a split-mouth design was chosen to monitor outcomes and possible discriminant factors. Here the synthetic material resulted more suited than silk as a suture as for the healing index and the patients' satisfaction. This may not seem completely in accordance with Banche et al. who compared silk to polyamide, polyesther and monocryl sutures finding silk as the least affine material to bacteria (13-33). It is to be underscored that nonresorbable sutures tend to accumulate more micro-organisms than the resorbable ones. On the other hand, a consistent outcome was described by Balamurugan et al. (34). These Authors tested silk suture and ab-



Healing Index: Healing Index is showed at 7 days.

sorbable polyglycolic acid suture clinically and histologically in 50 patients, after 7 days. They concluded that polyglycolic acid suture was superior, as "it had less tissue reaction, better handling characteristics and knotting capacity". When polyamide suture was compared with polyglactin suture (6) in oral implant surgery, no difference could be found between the two suture materials.

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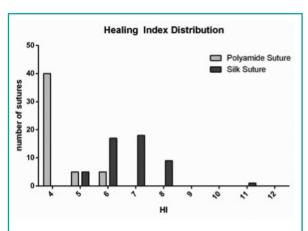


Figure 2
Healing Index distribution: Number of sutures in Polyamide or in Silk are showed for each Healing Index.

The present paper ads a clinical point of view to a previous *in vitro* report (35). Although there may not be only one ideal suture material the main factors to be taken into consideration are the surgery type and the tissue quality.

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