Ligature-Induced Peri-Implantitis in Minipigs Revisited

Abstract

Aim: The ligature-induced defect model still remains the model of first choice to experimentally investigate the cause, effect and treatment approaches of peri-implantitis. It was the aim of the present in-vivo trial to revisit the ligature-induced peri-implantitis minipig model regarding its current scientific value and ethical justification in implant research.

Materials and methods: Six minipigs were used for the analysis of peri-implant hard and soft tissue structures. Animals were randomly allocated to an experimental silk ligature-induced peri-implantitis group (n=4 animals) and a reference healthy group (n=2 animals). After six weeks mean pocket depths (PD) and bleeding on probing (BOP) measurements were performed just before animals were sacrificed.

Results: Overall, ligature-induced peri-implantitis provoked a local inflammation around the experimental implants. Additionally, a loss of crestal bone surrounding the implants could be detected. Mean pocket depths (PD) were 2.2 ± 1.1 mm for healthy animals and 5.4 ± 1.9 mm for peri-implantitis sites. Healthy sites showed a BOP of 60%, whereas peri-implantitis sites disclosed a BOP of 90% within 10 s after probing.

Conclusion: Clinical, radiological and histological findings of the present animal experiment supported the overall applicability of the ligature-induced peri-implantitis minipig model. A rapid breakdown of peri-implant hard tissues could be detected mainly on the buccal side.

Keywords: Mucositis; Bone loss; Inflammation; Dental implant; Animal model

Introduction

Treatment with osseointegrated dental implants has become a predictable and sustainable therapy for functional and esthetic reconstruction after tooth loss [1]. Yet, even though osseointegration of dental implants has clinically, anatomically, histologically, and ultrastructurally proven successful on long-term results inflammatory soft tissue issues, and there, mainly peri-implant mucositis and peri-implantitis are emerging problems [2-4]. In their consensus report Lindhe et al. concluded that peri-implant mucositis occurred in 80% of subjects and in 50% of implant sites, whereas peri-implantitis was identified in 28% and >56% of subjects and in 12% and 43% of implant sites [5]. This renders peri-implantitis a true threat in today’s advancement of implant reliability and performance.

In assessing the pathogenesis of peri-implant diseases, experimentally ligature-induced peri-implantitis lesions in animals have demonstrated histopathological similarities to naturally occurring lesions in humans [6]. Even though this chronic-type defect model using peri-implant ligatures was...
introduced in different large animal models, canines have established themselves as the most preferred species in the last decades. Schwarz et al. see the most distinct benefit of applying the canine model in the comfortable manageability by facilitating postoperative oral hygiene [7]. Furthermore, as anatomical dimensions of jawbones allow for placing conventional implants the authors regard canines as “[…] the most suitable animal species to conduct the ligature-induced peri-implantitis defect model […].”

According to Martini et al. however, the known preference of applying dogs in research was mainly based on the fact that knowledge, scientific findings and instrumentation from veterinary therapeutic interventions could be easily transferred to the needs of in vivo testing [8]. This finally led to the widespread use and acceptance of this animal species, even if other large animal models were available and the use of companion animals was highly controversial. Being in agreement with this perception it was the aim of the present in-vivo trial to revisit the ligature-induced peri-implantitis minipig model regarding its current scientific value and ethical justification in implant research [9]. The hypothesis of this experimental study in mini pig was that ligature-induced peri-implantitis causes a hard tissue breakdown at the buccal site. The stated null hypothesis (H0) was that there are no differences in pocket depth (PD) and bleeding on probing (BOP) between healthy and infected sites.

Materials and Methods

Animals and surgical model

Six neutered male minipigs (Ellegaard Göttingen Minipigs A/S, Dalmose, Denmark) were used for this study (age: 1.4 ± 0.0 years; weight: 34.25 ± 1.6 kg). Animals were randomly allocated to an experimental silk ligature-induced peri-implantitis group (n=4 animals) and a reference healthy group (n=2 animals). After extraction of the premolars and a healing period of 8 weeks, each n= 4 dental implants (SPI® Element Inicell®, PF 4.0, length 8.0 mm, Thommen Medical AG, Grenchen, Switzerland) were placed in each mandible. After six weeks animals were sacrificed. All experiments were conducted according to the Swiss laws of animal protection and welfare and were authorized by the local federal authorities (authorization #73/2013).

Animal care

Minipigs were acclimatized at least 30 days prior to surgeries to the new husbandry, bacterial environment and feeding. Clinically, a veterinarian examined animals 1-2 days after arrival and before surgeries. Twice daily, animals were scored (alertness, posture, appetite, respiration, signs of pain, lameness, temperature) by a veterinarian and a trained veterinary technician. In patient records were kept for each animal beginning the day of arrival documenting daily observations, treatments, surgical protocols and postoperative recovery. If abnormal findings were found, a senior veterinarian was immediately notified. Only healthy animals without any visible signs of illness were used. Upon arrival, feed was transitioned during a 3 week period from the supplier’s feed to pre-soaked, soft feed (600 g per animal per day, fed twice daily; KLIBA NAFAG 3000, Provimil Kliba AG, Kaiseraugst, Switzerland). Apple slices, apple sauce or yoghurt was fed by hand daily to tame the animals and was also used to give oral medication postoperatively. Enrichment included straw as bedding, toys, as well as a chain and wood to chew on. Even though animals were neutered, they showed excessive dominant behaviour when kept in groups of two or three. Animals were therefore kept solitary during the duration of the study but with visual contact to each other. Animals were regularly weighed at 2-4 week intervals. After surgery, enrichment was reduced to straw as bedding.

Anaesthesia and perioperative management

The day prior to surgery the animal was fed normally, bedding was removed overnight and feed was withheld preoperatively. Water was available ad libitum. After deep sedation in the stable (ketamine 20-40 mg/kg BW, Ketanarkon, Streuli Pharma AG, Uznach, Switzerland; midazolam 0.2 mg/kg BW; im, Sintetica S.A., Mendrisio, Switzerland), the animal was transported to surgery, an intravenous catheter was placed in an auricular vein, anesthesia was induced (propofol, to effect; Propofol® 1% Fresenius, Fresenius Kabi AG, Stans, Switzerland) and the animal was intubated in sternal recumbency. Carprofen (2 mg/kg BW iv; Rimadyl®, Pfizer AG, Zurich, Switzerland), penicillin (10’000 IU/kg BW iv; Penicillin Natrium Streuli®, Streuli Pharma AG, Uznach, Switzerland) and methadone (2 mg/kg BW iv; Methadon Streuli®, Streuli Pharma AG, Uznach, Switzerland) were given perioperatively. Right before surgery of each side, the respective mandibular nerve was locally anesthetized (ropivacaine, to effect; NAROPIN Inj Lös 0.5%, AstraZeneca AG, Zug, Switzerland) at its entry side into the mandible. The effect was controlled using a peripheral nerve stimulator (Innervator®, Fisher and Paykel Healthcare, Melbourne, Australia). Adrenaline in a sterile solution (adrenaline, diluted to 1:200’000; ADRENALIN Amino Inj Lös, Amino AG, Neuholw, Switzerland) was injected in the local mucosa to reduce bleeding. Intraoperatively monitored parameters included electrocardiogram, heart rate, pulse rate, arterial blood pressure (ventral tail artery) and oxygen saturation. Anesthesia was maintained via inhalation anesthesia (isoflurane in oxygen; Attane Isoflurane, MINRAD INC., Buffalo, NY, USA) and a CRI of propofol for a balanced protocol. Body temperature was supported through a heating mattress and/or a ventilation system. Blood was taken for routine hematology and blood chemistry analysis to provide a foundation for treatment in case of postoperative complications or prolonged recovery.

After surgery, animals were transported back to the stable accompanied by a veterinarian and closely observed until full recovery. Animals were covered during postoperative transportation and recovery to decrease loss of heat due to convection. Heating lamps and ventilation systems were available in the stable in case of hypothermia. Surgical sites were treated with cooling packs to reduce swelling. The IV catheter was removed the day after the surgery.

Pain medication (methadone 1-4 mg/kg BW iv; or buprenorphine 0.015 mg/kg BW im or iv; Temgesic®, Reckitt Benckiser AG, Wallisellen, Switzerland) was given postoperatively, antibiotics (amoxicillin 500 mg, Synulox®, Pfizer AG, Zurich, Switzerland) and anti-inflammatory drugs (carprofen 4 mg/kg BW po) were given for 5 days.
Tooth extraction

The animals were placed in lateral recumbency and the mouth was kept open with a mouth gag. The head was positioned using a moldable surgery cushion. Adhering to the surgical principle of adequate access, sulcular incisions were performed around the premolars. Additionally, a slight mesial vertical releasing incision was performed to allow a careful elevation of a full thickness flap. Standard dental instruments (forceps, elevators) were used to loosen and extract teeth. Crowns of the molars were vertically and horizontally separated (iChiropro, Bien-Air Dental SA, Biel, Switzerland). Root remnants were either removed with special root elevators or were drilled out with the same dental drill under adequate irrigation. Extraction sockets were cleaned. The lingual soft tissue was loosened from the bone plate and the mucoperiosteal flap was gently retracted with an elevator. Afterwards, the alveolar bone crest was down-leveled for 1 to 2 mm and sharp bony edges were smoothened.

The buccal and lingual mucoperiosteal flaps were repositioned and closed using single sutures (Vicryl® 2/0, Ethicon, New Jersey, NY, USA). The animal was then turned to the other side and the surgical procedure was repeated in an identical manner on the other side of the mandible.

Implant placement

Following eight weeks of healing the anaesthetized mini pigs were placed analogous to the procedure of the first stage surgery. The alveolar ridge was accessed through a full-thickness flap using a slightly lingual, mid-crestal incision in combination with slightly curved vertical releasing-incisions at its mesial and distal end. The flap was then elevated and held back using special retraction hooks. Implant sites were prepared according to a standard and approved drilling protocol using rotating pilot and twist drills in ascending order (diameter). After careful removal of bone debris from the drill holes with sterile 0.9% physiological saline, implants were inserted automatically (iChiropro, Bien-Air Dental SA, Biel, Switzerland). Root remnants were either removed with special root elevators or were drilled out with the same dental drill under adequate irrigation. Extraction sockets were cleaned. The lingual soft tissue was loosened from the bone plate and the mucoperiosteal flap was gently retracted with an elevator. Afterwards, the alveolar bone crest was down-leveled for 1 to 2 mm and sharp bony edges were smoothened.

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In the four animals of the experimental group, a single silk ligature (4.0) was placed around the abutment and slightly pushed downwards into the pocket. This finally led to a monolayer partly “submucosal” application. Ligatures were checked and maintained after two and four weeks (Figure 2). To reduce the burden of the animals and follow the 3R principles PD and BOP measurements were only performed at 6 weeks (time of sacrifice). BOP was evaluated as present if bleeding was evident within 10 s after probing, or absent, if no bleeding was noticed within 10 s after probing. PD was measured from the mucosal margin to the bottom of the probeable pocket. Each two measurements (mesial and distal) were performed at the lingual and buccal site. The statistical analysis was performed using a commercially available software program (SPSS® 18.0, SPSS Inc.). Mean values of all parameters were calculated.

Results

In the present study ligature-induced peri-implantitis provoked a local inflammation and loss of crestal bone surrounding the implants in four out of four animals. Thereby, probing depth, and the presence of bleeding characterized a soft tissue breakdown, whereas microradiographs and histological sections confirmed a peri-implant bone loss with vestibular dehiscence defects (Figures 3 and 4). For healthy animals the mean pocket depths (PD) was 2.2 ± 1.1 mm. In contrast mean pocket depths for peri-implantitis sites were 5.4 ± 1.9 mm. Healthy sites showed no bleeding within 10 s after probing (60%) and a slight bleeding in 40% of the cases. In contrast 90% of peri-implantitis sites disclosed a severe bleeding within 10 s after probing. Only 10% of sites showed no bleeding after 10 s. Clinically, bleeding on probing and a pocket depth of more than 5 mm demonstrated a diseased status after 6 weeks in the ligature-induced peri-implantitis animals. Furthermore, the peri-implant gingiva was slightly swollen and disclosed an intensive reddening. In contrast healthy animals showed some plaque formation and food remnants around the implants respectively the gingival formers. Yet bleeding was less conspicuous and peri-implant gingiva was less affected.

Overall, decrease of bone height was more prominent on the buccal aspect in all animals of the experimental group. Histologically, the measurable loss was up to three implant threads from bone level. On the lingual aspect the bony atrophy in the vertical direction was less distinctive with a mean bone loss of one implant thread. In half of the cases a formation of
a bony pocket respectively crater could be detected. Mostly, however, resorption was not limited to the outer diameter of the silk ligature, but was more comprehensive. Soft tissue structures disclosed a pronounced detachment in all animals of the peri-implantitis group, whereupon healthy minipigs revealed a tight sealing along the whole implant collar and gingival former (2.5 mm ± 3.2 mm).

**Discussion**

Until today, the ligature-induced defect model still remains the model of first choice to experimentally investigate the cause, effect and treatment approaches of peri-implantitis [10]. By using this model, however, Alhag et al. reported that several factors such as clot adhesion/stability and cellular migration/differentiation have to be carefully considered in the healing capacity of the peri-implant defect [11]. Therefore, Kolonidis et al. introduced an alternative model in Labrador dogs [12]. The authors placed titanium implants supracrestally allowing dental plaque to accumulate on exposed implant threads. After 5 weeks implants were cleaned and placed into fresh implant osteotomies on the contralateral side of the mandibles. Principally, this approach was based on the known perception that most dogs have a general tendency to develop a periodontal disease [13]. Even though the model was scientifically targeting to the right direction, burden and pain for the animals were ethically quite worth discussing. Beneath the much faster bone remodelling in dogs profound ethical constraints made this model do not become widely accepted. As similar emotional and legal issues count for the use of non-human primates in peri-implantitis research, their use is not well established in the European Community [14].

Generally, requirements and selection of a suitable animal model for peri-implantitis include the necessity to investigate the pathogenesis of the disease as well as the host’s potential and degree of regeneration. Concerning overall bone composition and bone remodeling, minipigs reveal a close similarity to humans and thus offer interesting aspects for analyzing osseointegration in dental and craniofacial research [15-17]. Microbiologically, Hickey et al. could also demonstrate that artificially induced peri-implantitis leads to a shift in the sulcular flora from primarily Gram-positive in healthy animals to Gram-negative in diseased animals [9]. Importantly, healthy animals do not disclose any specific species being characteristic for peri-implant lesions [7].

In conclusion, clinical, radiological and histological findings of the present animal experiment support the overall applicability of the ligature-induced peri-implantitis minipig model. The animals showed a progressive inflammatory soft tissue reaction with subsequent rapid breakdown of especially buccal peri-implant hard tissues. Operative prerequisites for generating the peri-implant lesions were silk ligatures and neglect of any plaque control. With respect to anatomical, physiological and ethical issues this model also proved to be a reliable and justifiable animal model. Further experiments have to elaborate and document the microbiologic and molecular pathways in this model for an improved insight into the natural development of the disease and thereby allowing a plausible translation to the human situation.

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