Surgical revision strategies for postoperative spinal implant infections (PSII)

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Abstract: Over the last years the number of spine surgeries with instrumentation has grown to an indispensable column in the treatment for different pathologies of the spine. A further increase in the incidence of instrumented spinal surgery is expected throughout the next years. Although the implementation and development of new techniques offer faster and more minimal invasive procedures, shortening surgery time, reducing soft tissue injury and revision due to hardware misplacement, the incidence of postoperative spinal implant infections (PSII) remains high. PSII related complications and revision procedures pose an enormous socioeconomic burden. Therefore, standardized strategies and protocols for treatment of PSII are urgently needed. While in former times hardware exchange or hardware removal was common practice in the field of spine surgery this approach has changed over the last years. Although the evidence from clinical studies in the field of PSII is of limited evidence, critical variables for revision strategies of PSII have been identified. Further, to quickly advance in the field of PSII it is certainly important to extrapolate and learn using data regarding the management from other fields of prosthetic joint infections. This should include clinical as well as experimental work in particular in the context of the biofilm, sonication as well as microbiological concepts. Over the last years, at our institution standardized procedures for diagnostic, surgical as well as antimicrobial treatment have been developed, based on the latest recommendations in peer-reviewed literature and our own data. Here we give an overview about surgical revision strategies for PSII and discuss the key points of our standardized protocol.

Keywords: Postoperative spinal implant infections (PSII); revision; biofilm; sonication; protocol

Introduction

The number of spine surgeries has been increasing over the last decades and spinal surgery with instrumentation has grown to an essential column in the treatment for various pathologies of the spine (1,2). Due to an ageing society, the rising use of spinal implants in the young as well as in the elderly, a further increase in the incidence of instrumented spinal surgery can be expected throughout the next years (1).

As demonstrated in single studies and confirmed by meta-analysis, the additional use of intraoperative imaging and navigation has helped to further improve pedicle screw accuracy as well as reducing invasiveness and thus soft tissue injury (3) in all regions of the spine, thereby potentially reducing revision rates and additional exposure to radiation and anesthesia (4-7). Although the implementation and development of these new techniques offer faster and more minimal invasive procedures the incidence of postoperative spinal implant infections (PSII) is reported from 1% up to 20% of all instrumented spinal procedures...
(8,9). Furthermore early PSII as well as chronic low virulent implant associated infections have been suggested to be associated with long-term hardware failure (10-14). Hardware failure often causes loss of spinal stability, resulting in pseudarthrosis, consecutive pain as well as recurrent spinal stenosis and back pain. Extensive revision surgery to replace the loosened screws or cages, along with extension of the construct and augmentation techniques, often have to be performed in the case of symptomatic loosened hardware (14,15). Finally every single case causes additional patient morbidity affecting long-term outcome, prolonging hospitalization and of course thus raising health care costs along with loss of working days (16). Therefore the management of surgical site infection following spinal instrumentation has become an important topic in the field of spine surgery (11,17,18). So far, the level of evidence from clinical studies in the field of PSII is very limited. However critical variables for revision strategies of PSII have been identified.

Since implantable devices are highly susceptible to bacterial colonization even low virulent bacteria can cause infection and recurrent infections due to biofilms, making them difficult to detect and eradicate (19,20). In former times hardware removal was common practise in the case of deep surgical site infection across different surgical fields using implants. In instrumented spine surgery especially implant removal is discussed ambiguously due to potential loss of correction even in fused patients (19,20). Over the last years, standardized procedures for diagnostic, surgical as well as antimicrobial treatment have been developed and implemented at our institution, based on the latest recommendations in peer-reviewed literature and our own data. Here we give an overview about surgical revision strategies for PSII and present the key points of our protocol as well as important general surgical aspects.

**Current evidence**

Despite of its increasing enormous clinical and socioeconomic importance (please see above), the level of evidence regarding the best treatment strategy of PSII is quite limited. So far no data from randomized controlled trials or prospective cohort studies are available in this field (8,9,11,18,23). However, the interest and importance of the topic is evolving, and lately a number of review and overview articles analysing the clinical data available have been published (8,11,15,18,24). However, it should be noted, that since the biofilm concept, advanced microbiological techniques (e.g., Sonication, PCR), the distinction between late and early infection along with an improved understanding of the pathogenesis of infected hardware as well as new hardware and surgical techniques have evolved significantly within the last years, data from retrospective single center cohorts analysing data, collected in part over more than 10 years (and reaching back to the 90’s) are hardly comparable. Therefore beyond this data, it is certainly important to extrapolate and learn using data regarding the management from other fields of prosthetic joint infections (PJI of the hip and knee) to quickly advance in the field of spine surgery (11,20,25-27). This should incorporate clinical as well as experimental work especially in the context of biofilm and microbiological concepts, too (19,20,28-30).

**Classification of PSII**

It is essential classify PSII. PSII should be classified as early, delayed or late infections. This classification is important for determining the most adequate treatment regime (see Figure 1). Furthermore, specific characteristics regarding the most probable pathogen and course may be derived from the respective subgroup (25,31). The classification of PSII and the respective treatment regime has been extrapolated and modified from other implant-associated infections (20,25,32).

Early infections are defined as infections occurring within 6 weeks after spinal surgery with instrumentation. Patients present most notably with acute local symptoms of infection: swelling, erythema, warmth, persisting surgical site drainage and/or fistula as well as possible systemic sings of infection like fever, increase in CRP, leukocyte cell count or erythrocyte sedimentation rate (ESR). However, especially laboratory results may be misleading as they have a low sensitivity especially with respect to low virulent pathogens, delayed PSII or in patients already receiving antibiotics (10,26,33,34). Delayed and late PSII occur more than 6 weeks after spine surgery (delayed within 1 year, late, defined as more than 1 year after surgery). Patients present with chronic wound drainage or fistula. Persisting or recurrent pain due to hardware loosing might be present as well. In delayed and late infections laboratory findings are often without pathological laboratory findings (33).

**Biofilm**

Implantable devices are highly susceptible to bacterial colonization and even low numbers of bacteria can cause
Microorganisms adhere to the implant’s surface and form biofilms (19,35). In general all bacteria are able to form a biofilm. In contrast to planktonic organisms, sessile bacteria within a mature biofilm are protected from phagocytosis, as well as other host immune responses (36). Further, as biofilm associated bacteria show an altered phenotype regarding growth rate and gene transcription they cannot be sufficiently targeted by antibiotics (28,36,37). Thus, the biofilm protects the microorganisms from the host immune system and renders them tolerant to antimicrobial treatments. Moreover, the biofilm hampers detection of the causative pathogen. Using the method of sonication, microorganisms can be released from the implants’ surface and quantitatively and qualitatively be detected from the detached biofilm in the sonication fluid. Thus to optimize detection in biofilm-associated infections, sonication of removed devices and prolonged incubation of cultures has been recommended (25). Sensitivity and specificity of sonication fluid is significantly higher compared to standard tissue cultures. Recent data showed that sonication of neurosurgical devices as well as pedicle screws is associated with a significantly higher rate of bacterial growth than in conventional cultures (14,38-40).

While in early implant-associated infections only an immature biofilm is found, in the case of delayed and late implant associated infections a mature biofilm is present. Staphylococcus aureus or gram negative bacteria are predominantly found in early infection, while the most common isolated pathogens in delayed infection are coagulase-negative staphylococci and cutibacterium acnes (formerly called propionibacterium acnes) (14,23,40,41). Moreover, it should be noted that beyond single species biofilms, as well multispecies biofilms in implant associated infection may be present (39,42). The key points including:

### Figure 1 Classification and procedure for PSII

<table>
<thead>
<tr>
<th></th>
<th>Early infection</th>
<th>Delayed and late infection</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>time</strong></td>
<td>≤6 weeks</td>
<td>delayed: &gt;6 weeks after</td>
</tr>
<tr>
<td></td>
<td></td>
<td>implantation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>late: &gt;12 months after</td>
</tr>
<tr>
<td>Clinical presentation</td>
<td>acute</td>
<td>chronic</td>
</tr>
<tr>
<td></td>
<td><em>local signs of infection, wound drainage, fistula, pain, fever</em></td>
<td><em>Chronic impaired wound healing, fistula, hardware loosening</em></td>
</tr>
<tr>
<td>Suspected pathogen</td>
<td>high virulent</td>
<td>low virulent</td>
</tr>
<tr>
<td></td>
<td><em>staph. aureus, streptococcus, gram. neg. bacteria (E. coli, Klebsiella, pseudomonas aerog)</em></td>
<td><em>coagulase neg. staphylococci. (e.g. staph epidermidis) anaerobic bacteria e.g. propioni bac. acnes</em></td>
</tr>
<tr>
<td>Biofilm</td>
<td>mature</td>
<td>immature</td>
</tr>
<tr>
<td>Surgical Treatment</td>
<td>DAIR possible</td>
<td>Removal or exchange of the implant necessary</td>
</tr>
<tr>
<td></td>
<td>Anti-biofilm treatment</td>
<td>(one-stage exchange)</td>
</tr>
<tr>
<td></td>
<td>Sonication of removed hardware</td>
<td>Anti-biofilm treatment</td>
</tr>
<tr>
<td>Antimicrobial</td>
<td>• usually 2 weeks intravenous treatment</td>
<td>• usually 2 weeks intravenous treatment</td>
</tr>
<tr>
<td>treatment</td>
<td>• following 4-10 weeks oral treatment</td>
<td>• following 4-10 weeks oral treatment</td>
</tr>
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modified from Conen et al. 2017
classification of PSII, suspected pathogen and the recommended procedure are summarized in Figure 1.

**Hardware retention vs. hardware exchange**

The effects of irrigation, debridement and implant retention (DAIR) in PSII versus hardware exchange/removal have been studied in a number of retrospective studies. Review of this studies along with the biofilm concept supports aseptic irrigation and DAIR followed by i.v. antibiotics and prolonged parenteral antibiotics as the treatment of choice for early PSII (9,11). As in early PSII an immature biofilm is present, the pathogen can be eradicated sufficiently due to its planktonic nature. Regarding the specific antibiotic regime please see the respective chapter on antibiotics.

In the case of delayed or late chronic infections a mature biofilm is present and thus hardware exchange is recommended, since the biofilm is cannot be completely eradicated from the implant’s surface. With hardware exchange or removal success rates up to 100% have been shown, while with DAIR in delayed infections recurrence rates between 20% and 50% occur (21,43-46). One study even reported a recurrence of infection in all patients with implant retention in treatment of delayed PSII. In this study cure was finally achieved after implant removal (21). As mentioned above, most of the studies are retrospective cohort studies, limiting the level of evidence. Lately the authors of a 20-year single center experience underlined the importance of implant removal/exchange in delayed infection versus DAIR in early infection (18).

Regarding exchange or removal of the interbody cage in PSII the available data is limited as well. Of course the incidence of PSII in constructs including an intervertebral cage depends on a number of different factors: stand alone cage or fusion, long/short construct, posterior, anterior or lateral approach. All of these factors impact the duration of surgery, extent of soft tissue damage, blood loss, transfusion and anaesthesia time. These factors have been identified as important procedural risk factors for PSII (8,9). Overall, the PSII incidence of an anterior or posterolateral instrumentation without a cage is lower compared to constructs including cages and posterolateral fusion (47,48). In contrast to pedicle screws and rods, removing or exchanging the interbody cage in PSII is thought to be associated with a potentially higher procedural risk due to scar tissue and the proximity to neural structures (11). So far, as available data suggests and as recently stated by an international consensus meeting on implants in infection after spine surgery, in PSII the cage can maintain if no signs of loosening along with bone loss and no signs of osteomyelitis or epidural abscess are present (11,49).

Of course, as different surfaces vary regarding their susceptibility to biofilm forming bacteria different cage materials have to be considered. However, although data from experimental work, showing lower susceptibility of titanium cages, the laboratory setting cannot be generalized for the clinical setting, as surface characteristics might also be affected by postoperative hematoma or seroma, as well as mechanical and thermal manipulation during the surgery. A number of studies have reported lower infection rates using titanium cages compared to stainless steel cages (50). Cages made of Polyethyleneetherketone (PEEK) have been reported to be associated with a higher infection rate compared to titanium (51). On the other hand, no difference of PEEK vs. Titanium cages was reported analysing data regarding surgical treatment of primary spinal infection (i.e., spondylodiscitis) (52,53). Taken together, the evidence level of titanium vs. PEEK is limited. In summary, as recommended by Divi et al. (11) in delayed PSII, the cage can maintain if no signs of loosening, bone loss and no signs of osteomyelitis or epidural abscess are present. Thus patients with deep PSII treated by DAIR and exchange of pedicle screws/rods and antibiotics should be followed up closely. If infection persists, cage exchange/removal should be considered if necessary using a lateral or anterior approach.

In some cases with chronic, deep spinal wound infection repeated surgical interventions might be necessary. The application of vacuum assisted closure (VAC) has been demonstrated to be an useful tool as the negative pressure promotes angiogenesis, the development of granulation tissue and reduces the number of bacteria. As recently reported VAC can efficiently and safely be applied in PSII even when the dura is exposed (54). Finally, in severe cases with significant wound defects and reduced soft tissue to cover the instrumentation and close the wound, a joint management applying VAC as well as complex flaps together with the plastic surgeons might be essential for successful treatment of PSII (55,56).

**General aspects in diagnosis and treatment of PSII**

As spine surgeons perform the initial surgery, of course they are as well consulted first if PSII is suspected. Most often, diagnostic measures as well as the surgical interventions are performed before microbiologists or infectologists are involved in the case. Thus, a number of
important points should be considered in the management of these patients including intraoperative aspects. These key points are listed below.

- Classification of early vs. delayed PSII is indispensable for adequate management;
- Laboratory markers are insufficient to rule out PSII;
- In all patients with local signs of infection PSII has to be ruled out as soon as possible;
- Early infection (≤6 weeks) can be managed with DAIR followed by i.v. and oral antibiotics;
- In delayed/late PSII change of implants is recommended;
- If not loosened or displaced, the cage can maintain;
- All devitalized, loose or purulent material, including non incorporated bone graft should be removed until vital (bleeding) margins are obvious;
- More than 3 intraoperative tissue probes using sharp dissection should be obtained for microbiology;
- Tissue samples should be obtained, where signs of infection are most prominent;
- Microbiological samples should be obtained from the screw canal as well;
- Sonication of removed hardware is recommended;
- Antibiotics should never be started before probes for microbiology are obtained;
- Always put a drain;
- Wound closure should be performed with donati single stich suture.

A standardized interdisciplinary protocol

At our institution all patients suffering PSII are treated by an interdisciplinary team including surgeons, infectologists and microbiologists. Standardized procedures for diagnostic and surgical as well as antimicrobial treatment have been developed based on the latest recommendations in peer-reviewed literature and our own data (10,11,14,25,26,31). Protocols can be found in the PRO-IMPLANT Foundation Guidelines (https://www.pro-implant-foundation.org/). Beyond conventional microbiological methods, sonication of removed hardware in PSII has been implemented as a routine microbiological procedure at our institution. The key points of our protocol are listed in Figure 1.

Summary

PSII associated complications and revision procedures pose a tremendous socioeconomic burden. Based on the available data and the latest recommendations we have implemented an interdisciplinary protocol at our institution. The following points build the main columns of the protocol:

(I) Early diagnosis and classification of PSII; (II) in the case of early PSII—early debridement and instrument retention followed by i.v. and parenteral antibiotics; (III) in the case of delayed/late PSII—debridement and exchange of instrumentation followed by i.v. and parenteral antibiotics; (IV) sonication of removed hardware; (V) close follow up of PSII patients.

The effectiveness of the interdisciplinary standardised protocol presented in this article has been studied in a retrospective manner with respect to prosthetic joint infections (PJI) of the hip and knee, as well as in a prospective manner for infections after cranial neurosurgery. Sonication of removed hardware in PSII has been implemented as a routine microbiological procedure in PSII at our institution. The value of sonication in instrumented spinal surgery has been demonstrated (10,14,20), especially with respect to chronic, low virulent infections. Regarding PJI protocol the rate of recurrent infection was significantly reduced form 10.4% to 3.1%. Applying the cranial protocol an overall infection free survival rate of 87% (27,32) was found. We strongly believe that standardized strategies and protocols for treatment of PSII will lead to a better outcome and reduce its socioeconomic burden. Prospective randomized controlled studies are urgently needed to evaluate the optimal treatment strategy for PSII.

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Footnote

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