A novel knee prosthesis model of implant-related osteomyelitis in rats

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Background and purpose   There have been numerous reports of animal models of osteomyelitis. Very few of these have been prosthesis models that imitate human conditions. We have developed a new rat model of implant-related osteomyelitis that mimics human osteomyelitis, to investigate the pathology of infection after orthopedic implant surgery.

Methods   2 wild-type strains of *Staphylococcus aureus*, MN8 and UAMS-1, and their corresponding mutants that are unable to produce poly-N-acetyl glucosamine (PNAG) (ica::tet) were injected into the medullary canals of the femur and tibia at 3 different doses: 10², 10³, and > 10⁴ CFU/rat. We measured clinical signs, inflammatory markers, radiographic signs, histopathology, and bacteriology in the infected animals.

Results   An inoculum of at least 10⁴ cfu of either wild-type bacterial strain resulted in histological, bacteriological, and radiographic signs of osteomyelitis with loosening of the prosthesis. An inoculum of 10³ CFU gave signs of osteomyelitis but the prosthesis remained in situ. Bacterial inocula of 10² cfu gave no signs of osteolysis.

Interpretation   We have established a new knee prosthesis model that is suitable for reliable induction of experimental implant-associated osteomyelitis with the prosthesis in situ, using a small inoculum of *S. aureus*. At a dose of 10³ CFU/rat, bacteria unable to produce PNAG (ica::tet) had only minor defects in their virulence.

Infections associated with in-dwelling orthopedic devices can be difficult to cure without removing the device, and they are quite expensive to manage (Darouiche 2004). Usual infection-control measures, laminair air flow, and use of systemic antimicrobial prophylaxis have not completely eliminated orthopedic implant-related infections (Darouiche 2003). *Staphylococcus aureus* is the predominant bacterium associated with infected metal implants (Eron 1985). There is an urgent need to explore other approaches to enhance diagnosis, prevention and treatment of infection, and osteomyelitis—for which appropriate animal models are extremely useful (Monzón et al. 2002). The aim of the present study was to establish an implant model of acute osteomyelitis associated with metallic implants without any promoters of infection other than the implant itself.

Materials and methods

*S. aureus strains and bacterial challenge (Table 1)*

We used the MN8 strain of *S. aureus*, originally obtained from a patient with toxic shock syndrome-1 (TSST-1), and the genetically related UAMS-1 strain of *S. aureus*, a primary isolate from a patient admitted to the Arkansas Children’s Hospital with an osteoarticular and soft-tissue infection and septic shock (Lucke et al. 2003, Cassat et al. 2006). UAMS-1 is a clinical isolate from osteomyelitis and is therefore clearly relevant to the model. Strain MN8 is closely related to UAMS-1 genetically: both belong to the EMRSA-16 group of *S. aureus*, which are characterized by an identical mutation in the alpha-toxin gene that results in loss of production of this factor. *ica*-deficient strains were used to evaluate the role of PNAG antigen, which is synthesized by proteins encoded at the ica locus (Cramton et al. 1999), to determine whether it is a dominant feature of implant-related infections caused by *S. aureus*. To date, 100% of over 100 clinical isolates from all sources tested have been found to express poly-N-acetyl glucosamine (PNAG), which contributes to this organism’s ability to make biofilms. Biofilms are thought to be a major component of the pathogenesis of implant-related infections.
Thus, PNAG may be a virulence factor in implant infections and is a known target for immunotherapy and vaccination.

41 rats were infected with 2 wild-type strains of *S. aureus* MN8 or UAMS-1, both of which are ica-positive, using inoculating doses of bacteria of $10^5$, $10^4$, $10^3$, or $10^2$ CFU/rat (MN8: $n = 23$; UAMS-1: $n = 18$). There were at least 3 rats in each challenge group.

Viable counts were done on the different bacterial strains to determine the exact dose inoculated. Both strains were also tested using mutants with deleted ica genes (ica::tet), which are referred to as ica-negative strains, at an infecting dose of $10^3$ CFU/rat with four rats in each group (Cramton et al. 1999, Beenken et al. 2004, Kropec et al. 2005). 5 rats were followed without being infected (the control group). The animals were followed for 2–6 weeks and then killed.

**Anesthesia**

All rats were sedated with a subcutaneous injection of hypnorm/dormicum, 0.3 mL/100 g body weight, given preoperatively and reinjected every 15 min (0.15 mL/100 g). After the operation, a femoralis block of the operated extremity was placed below the inguinal ligament using 1% lidocain/0.5% bupivacain in 1 mL. The rats were killed with an intracardiac injection of 2 mL pentobarbital (200 mg/mL).

**Surgery**

We used 54 Sprague-Dawley male rats that were 7–9 weeks-old (Taconic Europe) and weighing about 300 g. The experiments were approved by the Animal Committee of Denmark (reg. no. 2005/561-1049 from September 19, 2005).

The skin over the left knee was sterilized twice with alcohol. The fur was removed with a hair razor. The knee was opened with a parapatellar medial incision and the tendon with the patella was dislocated laterally. The articulating cartilage was osteotomized with bone scissors from the distal femur, with the proximal tibia including the menisci and cruciate ligaments protecting the collateral ligaments. A 2-mm-wide and a 10-mm-deep hole was bored with a hand drill into the femur and tibia to fit the joint components. The joint capsule and skin were closed with Ethibond 4-0 and Vicryl 5-0 after placement of the knee prosthesis (press-fit model) without bone cement.

Table 1. Inoculum, ica:tet status, and strain

<table>
<thead>
<tr>
<th>Group</th>
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<th>Inoculum (CFU)</th>
<th>ica status</th>
<th>Strain</th>
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<tr>
<td>Control</td>
<td>5</td>
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<td>MN8</td>
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<td>G1</td>
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<td>G2</td>
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<td>G5</td>
<td>8</td>
<td>$10^2$</td>
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<td>G6</td>
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<td>UAMS-1</td>
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<td>G7</td>
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<td>$10^3$</td>
<td>$-$</td>
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<td>G8</td>
<td>9</td>
<td>$10^4$</td>
<td>$+$</td>
<td>UAMS-1</td>
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**Prosthesis**

A rat-sized, non-constrained knee prosthesis was used (Figure 1) The femoral component was made from a metal alloy and the tibial component was milled from high-density polyethylene stock.

**Infection process**

The infection was initiated by direct injection of 10 µL of the appropriate *S. aureus* bacterial suspension into the medullary canals of the femur and tibia. The volume of the suspension fitted into the marrow hole. Afterwards the condylar prosthesis was inserted.

**Radiographic evaluation**

AP and lateral radiographs were taken on days 0 (OP), 7, 14, 21, and 42. To assess development and progression of bone infection, a modified scoring system was used (Schmidmaier et al. 2006). The following 6 parameters were scored: (I) periosteal reaction, (II) osteolysis, (III) soft-tissue swelling, (IV) deformity, (V) general impression or destruction, (VI) loosening of the prosthesis, and (VII) sequestrum formation. The score for each of the first 5 parameters used a scale of 0 to 4: 0 (absent), 1 (mild), 2 (moderate), 3 (severe). Parameter VI “loosening of the prosthesis” was judged as 0 (none), 1 (1 component), or 2 (both components), and parameter VII was judged as 0 (absent) or 1 (present). The maximum score that could be achieved was 18 (Figures 2 and 3).

**Microbiological evaluation**

The prosthesis components were explanted and rolled over non-selective media (5% Danish blood agar and chocolate agar plates (State Serum Institute, Copenhagen, Denmark). Isolated bacteria were identified as previously described (Højby and Frederiksen 2000). The same procedure was used for bone from the femurs and tibias and for the synovialis, which were rolled over Danish blood agar and chocolate agar in 3 directions and incubated at 37°C. Bacterial growth was judged as no growth, growth in the first streak (+), growth in the first 2 streaks (++), or growth in all 3 streaks (+++).
Bone and soft-tissue histology (Figure 4)

After removal of the prosthesis, the remnants of the tibia, femur, and synovialis were fixed in 4% buffered paraformaldehyde and decalcified in 10% formic acid/EDTA for 7 days. Samples were embedded in paraffin and transverse sections of 5 µm, including the implantation site, were cut on a microtome. The sections were stained with hematoxylin and eosin. For the first 30 rats, histological analysis of tissues from the blood, liver, and spleen was also carried out.

Semiquantitative scoring of all specimens was performed blind by pathologists who were not aware of the treatment groups (SSP and BMN).

For histological scoring of severity of inflammation, transverse sections of the tibia and femur (with the prosthesis removed) and tissue from the synovialis were investigated. Each of the 3 tissues (femur, tibia, and synovialis) was given a score ranging from 0 to 4, depending on the severity of inflammation. 0 meant no signs of inflammation, 1 was slight focal accumulation of inflammatory cells (neutrophils), 2 was a moderate but consistent inflammation in the transverse sections and/or moderate inflammation of the entire circumference around the cavity after prosthesis removal, 3 was the start of formation of an abscess in the cavity, and 4 was abscess formation and destruction of bone material with the synovialis completely infiltrated by neutrophils. The scores from the 3 separate tissues were added, giving a maximum score of 12.

Clinical evaluation

Body weight and temperature were determined, and well-being of the animal was estimated using food intake as a parameter. Macroscopic judgement of the knee was scored as described previously (Petty et al. 1988). Briefly, normal = 0, moderate swelling and edema = 0.5, swelling and abscess = 1.0, and abscess and illness of the animal = 1.5. If abscesses and/or fistulae were present, pus was aspirated and cultured on blood agar plates, and any bacteria were identified (Høiby 2000).

Statistics

All results are expressed as means. They were analyzed using the Kruskall-Wallis test for one-way analysis of variance. Statistical calculations were performed using SPSS for Windows software version 12.2. All p-values less than 0.05 were considered to be statistically significant.

Results

Radiography

No osteolysis or signs of inflammation were seen in any of the radiographs in the control group, including loosening of the prosthesis (Figure 1). Radiographic changes in groups G1 and G5 (rats injected with $10^3$ bacteria) indicated slight inflammation with little osteolysis and no loosening of the prosthesis. The modified scores (Schmidmaier et al. 2006) after 1 and 2 weeks were 2.9 and 3.6, respectively. Radiographs from most rats in groups G2 and G6 (injected with $10^3$ bacteria, ica+) showed slight to moderate inflammation with osteolysis, a periosteal reaction, and soft-tissue swelling but the prosthesis remained in situ. The scores for G2 were 1.4 and 5.6 after 1 and 2 weeks, respectively, and for G6 they were 0 and 5.3 after 1 and 2 weeks, respectively. The rats infected with $10^3$ CFU of the 2 different ica+ S. aureus strains MN8 and UAMS-1

Biochemical analysis

α-1 acid glycoprotein (AGP)—an acute-phase protein—was measured on days 0 (preoperatively), 7, 14, 21, and 42 (normal range: 0–130) (Matsumoto et al. 2007). Hemoglobin and leucocytes were also measured. There were no differences in the groups before and after the operation, irrespective of the bacterial strain and the infectious dose used.
(groups G3 and G7) had the same level of virulence as judged from the same scores of 7.4 and 5.5 after 2 weeks, respectively. The radiographic images of most rats in groups G4 and G8 (injected with $>10^4$ S. aureus MN8 or UAMS-1, respectively) showed moderate to severe inflammation and most of the prostheses had loosened or were no longer in place ($n=12$). For G4, mean scores after 1 and 2 weeks were 5.0 and 7.1, respectively, and for G8 mean scores of 7.2 and 7.8 were measured after 1 and 2 weeks (Table 2).

**Microbiology**

No bacteria were cultured from the control group; thus, the prostheses were sterile. In G1 and G5 inoculated (with $10^2$ CFU S. aureus), only 2 of 6 cultures obtained from the knees yielded a few colonies of S. aureus and these were seen only with the UAMS-1 strain. In G2 (wild-type strain MN8, $10^3$ CFU) 8 of 12 implant cultures had a moderate number of colonies (+++) and in G6 (wild-type strain UAMS-1, $10^3$ CFU), all 6 cultures were positive (++). In G3 (ica- S. aureus MN8, $10^3$ CFU), 3 of 4 cultures were positive at the ++ level. In G7 (ica-S. aureus UAMS-1, 103 CFU), all 4 cultures yielded colonies at the ++ level. In G8 (wild-type MN8 and wild-type UAMS-1, >$10^4$ CFU), all cultures were positive for S. aureus at the +++ level. No bacteria could be cultured from spleen, liver, or blood.

**Histopathology**

The histopathology was consistent with the radiographic findings. No signs of inflammation were found in the control group. No histological signs of inflammation or bacteria were found in the liver, spleen, or blood in any of the groups. In the 2 groups that were given an inoculum of $10^2$ (G1 and G5), all histological sections showed minor bone inflammation around the prosthesis with little bone destruction. In G2 and G6 (wild-type MN8 and wild-type UAMS-1 strains), the histopathology sections showed typical signs of bone inflammation (osteomyelitis) such as destruction of cortical and cancellous bone (osteolysis) and new bone formation. In G3 and G7 (the S. aureus ica-groups) the same findings were obtained but with less osteolysis. In G4 and G8 (wild-type MN8 and wild-type UAMS-1, >$10^4$ CFU), there were massive signs of bone infection with destruction and abscess formation. 4 rats were killed or died preoperatively because of aspiration (Table 3). The 2 groups infected with more than $10^4$ CFU were treated as 1 group for further analysis.

**Biochemistry and blood leukocyte counts**

In the control group, the preoperative AGP levels were < 200 µg/mL. In G2 (wild-type MN8, $10^3$ CFU), the AGP increased from a mean of 170 µg/mL to 542 µg/mL after 2 weeks, and then decreased to a mean of 424 after 4 weeks. In G6 (wild-type UAMS-1, $10^3$ CFU), the AGP increased from a mean of 168 to 218 after 2 weeks. The rats cleared the infection after 2 weeks. Hemoglobin and leucocyte levels were not statistically significantly different from those in the control group, and the biochemistry parameters after 2 weeks and the leucocyte counts and hemoglobin levels were similar. Only the AGP parameter was relevant for measurement of inflammation.

**Clinical results**

There was no statistically significant difference in mean weight loss between the control group and the G1 to G3 (CFU ≤$10^3$ and G5 to G7 rats. However, the weight loss in G4 and G8 rats given the highest dose ($>10^4$ CFU) was higher than in the control group. There was no soft-tissue swelling around the knee in the groups given $10^2$ CFU (G1 and G5). Body temperature was similar in the different groups.

**Discussion**

Various experimental animal models of osteomyelitis have been developed since 1884 when Rodet and Lexer (Rodet 1885, Lexer 1894) demonstrated that osteomyelitis similar to
that seen in humans could be created experimentally by injection of staphylococci to produce bone abscesses. Mader (1985) and Norden (1988), together with several other authors, demonstrated that osteomyelitis could be induced in an implant model by injection of bacteria into the tibia and femur cavities using either rats or rabbits (Mader 1985, Rissing 1990, An et al. 2006). Osteomyelitis studies related to orthopedic implant and prosthetic joint infection have been conducted using various animals such as rats (Zak et al. 1982, Rissing et al. 1985), rabbits (Nelson et al. 1990), dogs (Fokushima et al. 2005), and guinea pigs (Ofluoglu et al. 2007). The in vivo animal models of osteomyelitis have certain advantages, which have provided us with a clearer understanding of the osteomyelitis process and treatments. The limitations of the models were mostly due to differences in the immune systems of the animals, the non-physiological way the bacteria were inoculated into the animals, the short observation periods, and the high bacterial doses used (Mader 1985). Rats are the second most widely used animal for these studies, which is why we also used them. In addition, it is simple to evaluate the histological and microbiological bone processes in rats. Development of the surgical procedure and use of loop-glasses and a specially designed small non-constrained knee prosthesis allows evaluation of the osteomyelitis process in spite of the small size of the animal.

In most osteomyelitis models, lesions are induced by sclerosing agents, arachidonic acid, and foreign bodies containing bacteria to facilitate bone infection. Nelson et al. (1990) and Fukushima et al. (2005) did not use sclerosing agents and foreign bodies in their rat osteomyelitis models. They infected the animals with a combination of Pseudomonas aeruginosa and S. aureus. Fukushima et al. (2005) examined the relationship between the inoculating dose of bacteria and the progression of osteomyelitis, and found that development of significant histological and radiographic signs of osteomyelitis required inocula of at least 6 \times 10^3 CFU. Other authors have used foreign-body implants such as stainless steel pins and K-wires implanted into the bone marrow together with the bacteria to study the osteomyelitis process (Hudetz et al. 2008, Li et al. 2008).

In our rat model the foreign body was implanted as a knee prosthesis, to mimic the human condition in which the osteomyelitis process is initiated in the bone marrow around the prosthesis. Osteomyelitis was seen after 1 week; it was localized to the primary infected area without spreading to the liver, spleen, or blood. We found that inoculation of at least 10^3 CFU of 2 wild-type S. aureus isolates resulted in significant histological, bacteriological, and radiographic signs of osteomyelitis with loosening of the prostheses. Smeltzer et al. (1997) found that development of radiographic and histologic signs of osteomyelitis required inocula of at least 10^4 CFU. They used the same strain as we used (UAMS-1). We also found that the prosthesis remained in situ with an inoculum of 10^3 CFU, but the prosthesis was expelled after inoculation with > 10^4 CFU (for either S. aureus strain). For both MN8 and UAMS-1, 10^3 CFU was the optimal inoculum for study of the osteomyelitis process around the prostheses.

The ica gene is a major operon for expression of a potential virulence factor, a polysaccharide involved in staphylococcal biofilm formation (PNAG). However, the role of biofilms in staphylococcal bone infection is still unclear (Knobben et al. 2007), and they may not require the PNAG polysaccharide for formation. Both of the ica-strains gave rise to osteomyelitis that was no different to that achieved with the wild-type strains at the same size of inoculum (Table 2). Hudetz et al. (2008) found that the presence of ica genes had a strong effect on biofilm formation in vitro and a weak effect in vivo.

Overall, it appears that S. aureus causes persistent infection in a knee prosthesis implanted into rats, requiring a small number of bacteria. The tissue pathology and infection were independent of the presence of the ica genes and PNAG-dependent biofilm formation. We have also shown that after 2 weeks, the rats can clear the inflammation with doses between 10^2 and 10^4, indicating that it is not possible to study the course of inflammation for longer time. The immune system of the rat is quite different from the human immune system, which may explain their ability to clear the infection after injection of such relatively high doses. Even so, our study shows that a dose of 10^3 CFU of S. aureus would be a suitable experimental condition with which to study active and passive immunization against S. aureus in the same knee prosthesis model.

No competing interests declared.

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References


