Contamination during removal of cement in revision hip arthroplasty
A CADaver STUDY USING ULTRASOUND AND HIGH-SPEED CUTTERS
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Instruments used in surgery which rotate or vibrate at a high frequency can produce potentially contaminated aerosols. Such tools are in use in cemented hip revision arthroplasties. We aimed to measure the extent of the environmental and body contamination caused by an ultrasound device and a high-speed cutter.

On a human cadaver we carried out a complete surgical procedure including draping and simulated blood flow contaminated with Staphylococcus aureus (ATCC 12600). After cemented total hip arthroplasty, we undertook repeated extractions of cement using either an ultrasound device or a high-speed cutter. Surveillance cultures detected any environmental and body contamination of the surgical team.

Environmental contamination was present in an area of 6 x 8 m for both devices. The concentration of contamination was lower for the ultrasound device. Both the ultrasound and the high-speed cutter contaminated all members of the surgical team. The devices tested produced aerosols which covered the whole operating theatre and all personnel present during the procedure. In contaminated and infected patients, infectious agents may be present in these aerosols. We therefore recommend the introduction of effective measures to control infection and thorough disinfection of the operating theatre after such procedures.

Recent studies have shown that high-speed cutters in primary hip arthroplasty and spinal surgery can produce aerosols. These aerosols, possibly contaminated with bacterial, fungal or viral agents, are spread over the operating room and contaminate the environment and all personnel present during the surgical procedure. There is also a risk of infection through contact with mucous membranes or small wounds or by inhalation for both health-care workers and other patients.

In revision hip arthroplasty, different tools and high-speed cutters are used for removal of cement from the femoral cavity, particularly ultrasound devices, which vibrate at a high frequency. These devices also produce aerosols consisting of a mixture of irrigation solution, blood, tissue debris and body fluids.

Our aim therefore was to measure the environmental and body contamination caused by an ultrasound device and a hand-guided high-speed cutter used for removal of cement from the femoral cavity.

Materials and Methods

We carried out cemented arthroplasty on the left hip of a male human cadaver through an anterolateral transgluteal approach. With the cadaver supine, implantation and removal of a Lubinus SPII stem (LINK, Hamburg, Germany) left a typical cement mantle in the femoral cavity. A complete surgical procedure involved routine draping and the use of a 170 cm high drape barrier separating the operating table from the anaesthetist. All personnel present during the tests, which included an anaesthetist, a surgeon (sitting on the left side), an assistant (on the right side) and one scrub nurse (beside the cadaver’s legs on the left side) wore water-resistant sterile surgical gowns, gloves, caps and Fluidmasks (Tecnol, Fort Worth, Texas) covering the whole face. The operating room had no air-filtration system. The cement was extracted using either a high-speed cutting device with a 6 mm ball cutter (Ultra Power System; Zimmer, Warsaw, Indiana) or an ultrasound device (Ultradrive; Biomet, Warsaw, Indiana). Extraction with each device took 35 minutes. After each test, all draping and clothing were removed and replaced. We then repeated (three times for each device) the implantation and removal of the implant, thereby renewing the cement mantle.

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We placed two needles in vastus lateralis over the exposed femur in order to stimulate blood flow in the surrounding soft tissue. Another needle was introduced into the femoral cavity through a hole drilled after implantation in order to simulate intramedullary bleeding. The needles were connected to a 1.5 l reservoir containing a saline solution contaminated with *Staphylococcus aureus* (American Type Culture Collection, ATCC 12600). Determination of the final concentration of colony-forming units (CFUs) was after growth in tryptic soy broth (Merck, Darmstadt, Germany) under aerobic conditions at 37°C for 24 hours and then incubation on Columbia blood agar (Becton Dickinson, Franklin Lakes, New Jersey) at 37°C for 48 hours. The resulting concentration was $3.7 \times 10^4$ CFUs/ml. Both devices used sterile saline solutions in their integrated irrigation systems.

Contamination was assessed by placing 48 Petri dishes containing mannitol salt agar (Merck, Darmstadt, Germany) at the table height (100 cm) at even intervals over an area of 6 x 8 metres. The dishes were opened immediately before starting to remove the cement and closed five minutes after completion of the test in order to allow for complete settling of the aerosol. This gave a total exposure time of 40 min-

Fig. 1

Median CFU units in all sectors of the room for the detection of environmental contamination for a) the ultrasound device and b) the high-speed cutter with 95% CIs of medians.

Fig. 2

Median CFU counts for body contamination of the surgical time for a) the ultrasound device and b) the high-speed cutter with 95% CIs.
utes. We also obtained body and face contamination cultures from all team members. After incubation at 37°C for 48 hours any micro-organisms present on the Petri dishes were differentiated by morphological, physiological and serological criteria. For identification of *Staphylococcus aureus* we used both a commercial test kit (Pasteurex Staph Plus; Sanofi Pasteur Diagnostics, Chaska, Minnesota) and the classical tube coagulation test, calculating the median CFU counts from the repeated tests.

### Results

We detected environmental contamination throughout the room (6 x 8 m) after the use of both tools (Fig. 1). The ultrasound device showed a lower number of CFUs and no bacterial growth in some of the dishes, but there was growth in some of the most remote dishes at the borders of the room. The high-speed cutter contaminated all dishes with a high number of CFUs in all areas.

For body contamination the ultrasound device produced a lower number of CFUs in the surveillance cultures, but both devices contaminated all members of the surgical team to some extent (Fig. 2). The anaesthetist was least contaminated. The patient’s head, although protected by the drape, also showed contamination. The 95% confidence intervals (CI) demonstrated a significant difference between the two methods with the upper level produced by the ultrasound device being considerably less than the lower level produced by the high-speed cutter.

### Discussion

There are two distinct aspects to the risk of infection from an environment which is contaminated by aerosols. The first is the contamination of health-care workers in the operating room either directly through the aerosol cloud or by contact with contaminated instruments or surfaces. There are several reports of infection from bacterial agents such as *Staphylococcus aureus* and viral agents like hepatitis B and C and *Herpes simplex* from injuries with sharp and high-speed tools. There is also a risk of infection for team members through inhalation of aerosols contaminated with pathogens such as *Mycobacterium tuberculosis*, legionella, hepatitis B, varicella zoster, smallpox, influenza and *Staphylococcus aureus*.

The second is a risk of infection for patients operated on in the same room after such surgery or in contact with contaminated medical staff. Evidence for such a route of transmission has been reported by Isenberg et al for a single-source outbreak transmitted through a scrub nurse with positive cultures of *Candida tropicalis* on her fingertips and in her nasopharynx. Other bacterial and viral agents can survive on surfaces in the operating room and then be transmitted by way of medical personnel and instruments.

Both devices for removal of cement contaminated all individuals in the operating room to some extent. Sufficient personal protection, consisting of water-resistant surgical gowns with long sleeves, gloves and full-face protection with face shields should be mandatory during these operations. Protection should be worn by everyone in the room including the anaesthetist, not only throughout the procedure, but afterwards until the room has been disinfectected.

The aerosol produced by both devices covered an area of 6 x 8 m, the whole operating room workspace. The number of CFUs differed for the two devices, with the ultrasound device producing much lower numbers. This may have been due to the high temperatures of over 100°C which were recorded during the use of this device; some of the most remote dishes were, however, contaminated. Although a lower number of CFUs means a lower chance of infection, there remains a risk. The extent, concentration and risk of infection were higher with the hand-guided high-speed cutter.

We simulated blood flow and blood loss comparable to those of a revision hip arthroplasty as simulation directly through the vessels of a cadaver is not possible. While the simulation of blood flow which is produced by placing needles in the soft tissues surrounding the femur and into the femoral cavity may be a limitation of the model, we believe that our results are comparable to those of a real procedure. A concentration of 2 to 4 x 10^6 CFUs/ml in the fluid gave a realistic simulation of an infected patient, since such and even higher plasma levels of infectious agents are found in vivo. *Staphylococcus aureus* was chosen as the example micro-organism because it is easily detectable. The use of a human cadaver meant that the surgical procedure was realistic and all draping and clothing were those routinely used in the orthopaedic operating theatre.

We propose that effective measures of body protection for all personnel during revision arthroplasty should be introduced. Only a minimum number of people, all trained in the appropriate safety procedures, should be present in the operating room. Air filtration in the operating theatre may lead to improved safety of the personnel and the patient. We also recommend that patients with proven infections are operated on at the end of the day and that the operating theatre is efficiently disinfected after all removal of cement using ultrasound devices or high-speed cutters.

No benefits in any form have been received or will be received from a commercial party related directly or indirectly to the subject of this article.

### References


