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Attitudes, risk of infection and behaviours in the operating room (the ARIBO Project): a prospective, cross-sectional study

Gabriel Birgand,1,2,3 Christine Azevedo,4,5 Gaelle Toupet,3 Roger Pissard-Gibollet,5 Bruno Grandbastien,6 Eric Fleury,5,7 Jean-Christophe Lucet1,2,3

ABSTRACT

Introduction: Inappropriate staff behaviours can lead to environmental contamination in the operating room (OR) and subsequent surgical site infection (SSI). This study will focus on the continued assessment of OR staff behaviours using a motion tracking system and their impact on the SSI risk during surgical procedures.

Methods and analysis: This multicentre prospective cross-sectional study will include 10 ORs of cardiac and orthopaedic surgery in 12 healthcare facilities (HCFs). The staff behaviour will be assessed by an objective, continued and prolonged quantification of movements within the OR. A motion tracking system including eight optical cameras (VICON-Bonita) will record the movements of reflective markers placed on the surgical caps/hoods of each person entering the room. Different configurations of markers positioning will be used to distinguish between the staff category. Doors opening will be observed by means of wireless inertial sensors fixed on the doors and synchronised with the motion tracking system. We will collect information on the OR staff, surgical procedures and surgical environment characteristics. The behavioural data obtained will be compared (1) to the ‘best behaviour rules’ in the OR, pre-established using a Delphi method and (2) to surrogates of the infectious risk represented by microbiological air counts, particle counts, and a bacteriological sample of the wound at closing. Statistics will be performed using univariate and multivariate analysis to adjust on the aerolic and architectural characteristics of the OR. A multilevel model will allow including surgical specialty and HCFs effects. Through this study, we will develop an original approach using high technology tools associated to data processing techniques to evaluate ‘automatically’ the behavioural dynamics of the OR staff and their impact on the SSI risk.

Ethics and dissemination: Approval of the Institutional Review Board of Paris North Hospitals, Paris 7 University, AP-HP (no 11-113, 6 April 2012). The findings will be disseminated through peer-reviewed journals, and national and international conference presentations.

INTRODUCTION

Surgical site infection (SSI) is a major public health problem. It is the third most common healthcare-associated infection and contributes to 13–17% of all such infections.1 2 In
guidelines do not include specific recommendations regarding the best OR staff behaviour (except for wearing cap, mask and scrub and performing appropriate hand hygiene) to decrease the exogenous risk of SSI. Some surgical or infection control societies advise to control the OR traffic in order to decrease air contamination and wound colonisation. These measures include limiting door opening and restricting the movements and the number of persons in the OR. However, these recommendations are often vague and are based on expert advice only without robust scientific arguments.

At the beginning of the 1980s, Lidwell et al. evidenced a correlation between airborne contamination and wound contamination. Later, Tammelin et al. demonstrated that surgical wound contamination could originate from the staff skin flora. Staff skin shedding could lead to spreading microorganisms by the air with occurrence of wound contamination. In addition, the impact of the staff behaviour on SSI risks was assessed in several studies through the observation of the number of persons in the OR and the frequency of door opening. Overall, these studies were performed using classical audits with evaluation by human observers over short-time periods. This methodology does not allow the objective, continued and rigorous full collection of behaviour in the OR. Indeed, behaviours may change in the presence of an observer and observations may differ from one expert to another.

Research questions
To date, no study has described the global OR staff dynamics and behaviours during surgical intervention in the OR. This issue is challenged by the methodological concern of observational sessions by a direct observer. In consequence, the impact of the OR staff dynamics and behaviours during surgical intervention in the OR on the SSI risk has been partially assessed in previous studies. New technologies using motion capture systems appear as an alternative to obviate methodological issues. Automatic techniques based on sensors or motion capture allow the acquisition of objective data, with continued and prolonged periods of data collection.

Study objectives
The present study will focus on the assessment and description of OR staff behaviours and on its association with the SSI risk during surgical procedures. This study aims at describing and assessing the staff behaviours in the OR and their variability by recording staff displacements using a motion tracking system and door opening detection system. Data obtained will be compared to the best practices previously established by an expert panel during an earlier part of the study. A secondary objective was to correlate the staff behaviours with the SSI risk, approached with surrogates of SSI, such as OR air contamination and wound contamination at the end of the surgical procedure.
The main objective was to objectively measure the movements of surgical teams in the OR and to assess their adherence to the pre-established best practice criteria and their variability in a panel of ORs from two surgical specialties.

Secondary objectives are:
1. To assess correlations between movements of the OR personnel and the SSI risk, as approximated by surrogates of the infectious risk;
2. To assess the correlation between the particle count and the microbiological contamination in the OR air;
3. To describe the change in practices in the OR depending on the presence of a motion tracking system;
4. To assess the OR staff perception of their behaviour in the OR during an intervention to correlate with the actual data.

**METHODS AND ANALYSIS**

**Study design**

We propose an observational study based on the correlation between the data on OR behaviours obtained using new technology tools and (1) the ‘best behaviour rules’ established by an expert panel during an earlier part of the study and (2) surrogates of the infectious risk in the OR.

**Population and location of the study**

The study population will be formed of the OR personnel (surgeons, anaesthesiologists, nurses, nurse’s aide) and any other person likely to enter the OR during the surgical procedure. Among this population, behaviours will be analysed by an automatic system of motion capture. Volunteers to participate will be selected by contacting the heads of the surgical and anaesthesiology teams and the infection control practitioner from each selected healthcare facility (HCF).

Surgical specialties and procedures have been included according to the following criteria: cutaneous approach, clean contamination class (Altemeier’s class I), the frequency and the reproducibility of the procedure. On this basis, two specialties will be included: cardiac surgery with procedures requiring a full median sternotomy (planned coronary artery bypass grafting, valve repair or replacement surgery); and orthopaedic surgery for total hip and knee replacement.

**Strength evaluation**

The strength estimation for this comparative study depends on the variability of the staff behaviour between ORs. We hypothesise that the behaviour’s variability is lower between the OR personnel in a same HCF than between two different HCFs. Thus, we have chosen to include a panel of HCFs to take into account this potential variability. We will include interventions occurring during half-day periods. For each surgical ward involved in the study, one OR will be randomly selected. The inclusion of 20 different ORs (10 in each specialty) will generate data for 50 cardiac procedures (1 patient per day) and for 50–100 orthopaedic procedures. The final analysis will be performed on a total of 100–150 procedures.

We will perform an observational multicentre study including 12 HCFs: 7 University hospitals, 1 semiprivate and 4 private hospitals located in France. Among the 12 participating HCFs, 10 OR of cardiac surgery and 10 OR of orthopaedic surgery (7 public, 1 semiprivate and 2 private for both specialties) will be included in the study.

**Judgement criteria**

**Primary criteria**

**Motion capture**

The main judgement criteria will be the staff behaviour, as measured using his/her movements in the OR. A technology of motion capture based on a video tracking system will be adapted for the objective, continued and prolonged detection and the characterisation of movements in the OR. A network of eight video cameras (VICON-Bonita, Vicon, Los Angeles, USA) will be fixed upright to the wall by a suction system and linked by Ethernet cables to a hub. Data will be recorded on a laptop using the Vicon Tracker software (Vicon). Briefly, 68 LEDs situated on each camera produce an infrared light reflected by hemispherical markers and acquired by the optic. The detection of the same marker by different cameras allows its three-dimensional (3D) positioning.

The motion capture will be performed by a continuous tracking of reflective markers placed on the surgical caps/hoods of each person entering the OR. This system evaluates the movements of persons with a precision of 50 cm. Four types of marker combinations will be created to distinguish four different professional categories: surgeon, anaesthesiologist (including anaesthesiology nurse and extracorporeal circulation personnel), OR nurse and other. The markers’ positions are located in 3D by a method of spatial triangulation. The cartography of the OR including the situation of the table and doors will be performed at the system installation. Data recorded by the system will include the time and 3D position of the barycentre of each marker combination.

Doors opening will be collected by autonomous inertial sensors fixed on each door and synchronised to the motion tracking system. HiKoB FOX (HiKoB, Villeurbanne, France) is an autonomic system of wireless inertial sensors (tri-axial accelerometers, gyrometers and magnetometers). HiKoB FOX collects dynamic and door movements in real time. One sensor will be fixed on each door of the OR after a temporal synchronisation with the motion tracking system. The consistency between the position of persons in the OR and the doors traffic will be controlled through this synchronisation.

This device including the motion tracking system and the inertial sensors will stay during 1 week in the same OR to get people used to it and to take into account the
potential behavioural modifications due to the Hawthorne effect. Data acquisition will start at patient entry in the OR and will continue until patient exit. Door opening sensors will be kept for one additional week to evaluate the impact of the Hawthorne effect on behaviour by comparing the frequency of door opening during and after removal of the motion tracking system.

Best practices
The best practices in the OR have been established using a Delphi method. During this procedure, we asked a French college of experts including five surgeons, five anaesthesiologists, five OR nurses and five infection control physicians, to quote a preselected list of behaviours and practices potentially linked to an increased SSI risk. These experts are professionals involved in surgical activities and are working daily in the field of infection control and SSI prevention. Overall, 24 experts were contacted and 20 agreed to participate. Items were preselected using the national and international guidelines for SSI prevention and were then discussed by a working group composed of one surgeon, one anaesthesiologist and three infection control physicians, all specialised in quality and safety in the OR. We selected 14 preoperative, 35 operative and 1 postoperative items. We asked to quote the infectious risk on a Likert scale varying from 1 (no impact on the infectious risk) to 9 (high impact on the infectious risk). Additionally, we asked to quote 14 additional items possibly collected by the motion capture system. An item was selected if quotations were higher than 6 for more than 17 participants.

Finally, 11 variables were considered to significantly increase the SSI risk. These variables were selected after two quotation rounds. Moreover, four parameters were considered to be interesting to collect with the motion tracking system (box 1).

Secondary criteria

Air sample
Microbiological air counts will be measured using an impactor air sampler (Air-test Omega, LCB, La Salle France) at a flow rate of 100 L/min for 5 min (500 L) sampling on to Trypticase soy agar (BioMerieux, France), which will then be incubated for 4 days at 30°C. Air counts will be expressed as colony-forming units/m³. The air sampler will be positioned at the head of the patient. After each sample, the impactor will be disinfected. Samples will be performed at the incision, every 30 min during orthopaedic surgery, every hour for cardiac surgery and at wound closing. The time of samples will be synchronised on the motion tracking computer clock.

Particle count
The particle count (HandiLaz Mini, Particle Measuring Systems, Boulder, USA) will be performed using a photo-detection device continuously from incision to wound closing. A base level of particle count will be performed for each OR at the beginning of the day before the staff entry. The particle analyser will sample 1 min every 2 min throughout the surgical procedure at a rate of 0.0283 m³/min (1.0 ft³/min) and logged data at 1 min intervals to obtain the sample volumes of 0.0283 m³ (28.3 L) of air. Samples will be collected through a 100 cm length of the surgical wound at the patient head. Particles will be classified by diameter (d) in six-size ranges: 0.3 ≤d<5.0, 5.0 ≤d<10.0, 1.0 ≤d<3.0, 3.0 ≤d<5.0, 5.0 ≤d<10.0 and d≥10 mm. The count and particle size measurements will be recorded electronically by the particle analyser from the patient entry to the exit of the OR. The particle analyser will be synchronised on the motion tracking computer clock.

Wound sampling
A sample of the operating wound will be performed before closing. The sample method previously described by Tammelin et al., will use sterile pads of polyamide-polyester-viscose measuring 7.5×7.5 cm placed on the subcutaneous tissue and removed after being soaked by wound liquids (1 min). This sample will be performed before any antiseptic aspersion. Microorganisms will be extracted by vortexing the pads during 2 min in a phosphate buffer saline (PBS with

<table>
<thead>
<tr>
<th>Box 1</th>
<th>The final criteria selected by a college of experts using a Delphi procedure</th>
</tr>
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<tbody>
<tr>
<td><strong>Parameters linked to an increase risk of surgical site infection (SSI)</strong></td>
<td></td>
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<tr>
<td><strong>Preoperative period</strong></td>
<td></td>
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<tr>
<td>1. Patient skin preparation (hair removal and skin antisepsis).</td>
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<tr>
<td>2. Wearing of a surgical cap covering all hair surfaces by all persons in the OR during the surgical procedure.</td>
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<tr>
<td>3. Wearing of a mask covering the nose and mouth by all persons in the OR during the surgical procedure.</td>
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<tr>
<td><strong>Operative period</strong></td>
<td></td>
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<tr>
<td>4. Frequent hand disinfection of the surgical team.</td>
<td></td>
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<tr>
<td>5. Quality of air ventilation (type of flux, type of air contamination, pressures).</td>
<td></td>
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<tr>
<td><strong>Postoperative period</strong></td>
<td></td>
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<tr>
<td><strong>Secondary criteria</strong></td>
<td></td>
</tr>
<tr>
<td>1. The knowledge and the actual implementation of the guidelines for SSI prevention.</td>
<td></td>
</tr>
<tr>
<td>2. Patient skin preparation (hair removal and skin antisepsis).</td>
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<tr>
<td>3. Setting up of sterile drapes.</td>
<td></td>
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<tr>
<td>4. Surgical hand disinfection of the surgical team.</td>
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<tr>
<td><strong>Operative period</strong></td>
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<tr>
<td>6. Permanent wearing of scrub suits by everyone in the operating room (OR) (mask/surgical caps).</td>
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<tr>
<td>7. Permanent wearing of specific sterile suits for the operating staff.</td>
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<tr>
<td>8. Wearing of a surgical cap covering all hair surfaces by all persons in the OR during the surgical procedure.</td>
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</tr>
<tr>
<td>10. Systematic replacement of material of wear in case of asepsis fault.</td>
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**Secondary criteria**

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Tween 80 at 2% and lecithin at 0.3%, Hyphen BioMed, Neuville sur Oise, France) inactivating antiseptics compounds. For each pad, an aliquot of 0.5 mL of PBS will be cultured on blood agar. Strains isolated will be quantified and identified by the investigator team using a mass spectrometry assay (MALDI-TOF-MS system, Microflex, Bruker Daltonics, Bremen, Germany).9

Data collection

Information will be collected on: (1) OR staff present during the intervention (age, gender, function, experience in the function, role in the surgical procedure), (2) surgical procedure (surgical specialty, surgical procedure, surgical technique used, incision time, preselected procedure periods, closure time), (3) surgical environment characteristics (air changes of filtered air per hour, positive pressure, temperature, relative humidity, particles contamination class, kinetic of particle decontamination class). The architecture of the OR will be taken into account by collection of the sizes and volumes of the room.

The motion tracking system will record all the behavioural parameters including: the number of persons in the OR, their proximity to the surgical theatre, the number of door openings, the number of exit/entry, the cumulated time in the OR and the interaction between the OR staff. These data will be stratified according to the professional categories.

As described above, particles’ contamination will be continuously collected at the head of the patient from entry to exit from the OR. Microbiological data will include qualitative and quantitative values of the total air and wound flora.

Additionally, the perception of each OR staff will be collected using two different questionnaires: one assessing the safety climate, previously used by Sexton et al.21 and the other assessing the perception of each person regarding the infectious risk in his/her OR (figure 1). These questionnaires will be given to every OR team member and will assess six dimensions: perception of management, safety climate, stress recognition, job satisfaction, working conditions and teamwork climate. Participants will give his/her perception by rating 59 items corresponding to these six dimensions on a Likert scale going from A (disagree strongly) to E (agree strongly). The results will be stratified by the professional categories.

Statistical analysis

A descriptive analysis of all the parameters collected will be performed. For continuous variables (ie, age, duration of presence in the OR), indicators such as the mean, SD, minimum, median, quartiles and maximum values will be calculated.

χ2 Test will be used to evaluate the homogeneity between the observed behavioural data and the items of interest previously established by the college of experts, both categorical variables, with a level of significance for

the p value fixed at <0.05. These analyses will be performed after stratification according to the surgical specialty and the type of HCF. Pearson or Spearman tests will be used to assess the correlation between the quantitative variables.

Additionally, a logistic regression will be used to independently analyse the link between the environmental variables (air wound contamination) and the behaviour variables all dichotomised according to the distribution of the population. The model will be adjusted according to the aerolic and architectural characteristics of the OR. The model will also be adjusted according to the Hawthorne effect with the comparison of door opening data obtained with and without the video tracking system.

Finally, a multilevel model will allow including contextual variation due to the surgical specialty or the type of HCF. The method will quantify the ‘surgical specialty’ effect and the ‘HCF type’ effect.

Confidentiality issue

The motion capture system will not allow identifying people who are symbolised by markers on the head cap. Functions of the OR team members will be collected but no name data will be recorded. The system will be presented to surgical and anaesthesiology teams in each participating centre and OR. Included patients will systematically be informed by an information letter. Additionally, this methodology requires the consent of the OR members included in the study.

DISCUSSION

The operating theatre is a particular area in the hospital. This special care environment with sophisticated techniques generates several ranges of risks for the patient including the occurrence of infection. Several recommendations have been published (skin preparation, surgical antibiotic prophylaxis, control of the OR environment and improvements in the surgical technique) to improve the patient safety and quality of care in the OR.10 11 24 25 Most of the recommendations are based on the scientific evidence. However, guidelines for the prevention of transmission of microorganisms to the surgical wound and eventually to SSI are scarce and often fuzzy. The existing recommendations are based on expert advice only without any scientific proof of evidence. In consequence, rituals abound in operating departments to prevent environmental contamination and impact staff behaviour. The present study will aim to bring a rationale for the prevention of airborne microorganism transmission by the description of best behaviour rules in the OR.

The impact of behaviours on the SSI risk has been studied in the past. We performed a search on the MEDLINE database in March 2013 and found eight articles assessing the correlation between the behaviour and the infectious risk in the OR. Results were conflicting. Among those, five articles studied the traffic flow by
evaluating the number of door openings and reason for door opening and the number of persons attending the OR during the intervention. Four of them were only descriptive studies and the last study correlated the traffic with the air contamination as a surrogate of the SSI risk.\textsuperscript{18, 19, 26–28} These studies were based on human observations performed at a given time. Collecting information using observers may have induced two biases: the Hawthorne effect and the partial observation of dynamics in the OR.

In two other articles, a bundle of preventive measures (including behavioural measures) were implemented and showed some impact on the SSI rates.\textsuperscript{8, 29} Finally, a study evaluated the impact of noise and found a strong positive correlation between the increase in decibels and the SSI rates.\textsuperscript{30} These studies were based on human observations of door opening or person present in the OR at given times. However, no study systematically and continuously evaluated a global perception of the staff dynamics in the OR.

Through the present study, we will develop an original approach using high technology tools (motion tracking and inertial sensors) to evaluate the behavioural dynamics of healthcare workers in the OR and their impact on the SSI risk. Other studies have used video to audit practices in the OR.\textsuperscript{31} This system was generally used to analyse and improve the performance of surgical techniques and to prevent an adverse event in the OR.\textsuperscript{32, 33} This type of recording system has also been used to improve practices outside the OR. A study showed that compliance with hand hygiene could increase from less than 10–86% due to the presence of remote video auditing.\textsuperscript{20}

This challenging project gathers specialists from several disciplines (infection control, epidemiology, surgery, anaesthesiology, psychology and engineering) and will allow obtaining the qualitative and quantitative epidemiological data. This consistent approach will allow collecting data that will help to understand the behavioural origin of the SSI risk and to improve the quality of care in the OR.

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Contributors CA, BG, GT, RP-G, EF, J-CL and GB have been involved in drafting the manuscript and have made contributions to the conception and design of the study.

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Competing interests None.

Ethics approval This protocol was approved by the Institutional Review Board of the (RB) of Paris North Hospitals, Paris 7 University, AP-HP (no 11-113, 6 April 2012). The findings of the study will be disseminated through peer-reviewed journals, national and international conference presentations.

Provenance and peer review Not commissioned; internally peer reviewed.

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