

## Editorial

# Progress in Surgical-Site Infection Surveillance

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How many surgical-site infections (SSIs) occurred last year in the United States? Were there more or fewer than 5 years ago? If more, could we have done better? If fewer, could we have done better still?

It has been more than 25 years since the Centers for Disease Control and Prevention's landmark Study on the Efficacy of Nosocomial Infection Control (SENIC) provided our first truly national portrait of surveillance for SSIs.<sup>1</sup> Among SENIC's contributions was a robust risk index whose direct descendant we continue to use.<sup>2</sup> A generation later, one would expect SSI surveillance methods to be well established, the epidemiology of SSIs to be well understood, and the research frontiers to be elsewhere.

And yet, in this issue of *Infection Control and Hospital Epidemiology* new reports from three continents speak to important unmet needs.<sup>3-5</sup> Two of these articles address postdischarge surveillance, a problem that increasingly claims our attention as the duration of postoperative hospitalization shrinks, often to zero. A consensus statement in 1992<sup>6</sup> asserted there was no validated method for conducting postdischarge surveillance, and this conclusion remains true.

The current report by Avato and Lai joins those of many predecessors<sup>7-22</sup> in noting that most infections first become manifest after discharge. This was true even though they studied coronary artery bypass graft surgery, a procedure whose postoperative length of stay is still measured in days rather than hours. This report illustrates the special circumstances currently required to establish an effective system, and it also indicates the magnitude of the effort. First, Avato and Lai were able to learn what the surgeons knew because they could establish ongoing coordination with the surgeons' offices. That, in turn, was possible because these offices were located in or near the hospital, allowing ongoing com-

munication and review of office-based medical records as necessary. Second, the hospital's infection control department was able to make an ongoing commitment of the scarcest and most valuable of its resources, a large amount of practitioner time. If the University of Massachusetts experience is typical, then performing postdischarge surveillance for all coronary artery bypass surgery in the United States will absorb the full-time effort of several hundred infection control practitioners. Comparable effort for all surgical procedures will require considerably more than the full-time effort of all current infection control professionals.

Thibon and colleagues remind us that the period of postdischarge risk often lasts longer than does postoperative care. Fortunately, the biology of SSI allows us to focus on the first 30 days for procedures that don't involve implanted prosthetic material. However, even 30 days is long, compared with the timing of follow-up office visits. So, we will need to incorporate the "when" of postdischarge surveillance into our methods and our lexicon. Survival analysis, the method adopted by the authors, is a good solution—if, as they point out, patients whose last follow-up occurs before 30 days are neither more nor less likely to experience infection during the remainder of the 30 days than patients whose last follow-up occurs at least 30 days after surgery. Is this true? We don't know. Is it a reasonable surmise? Possibly. In any event, the authors make a strong case for describing the duration of follow-up in any report of postdischarge surveillance.

The article by Russo and Spelman addresses the fact that all patients are not created equal regarding their baseline risk of infection. Our best case-mix adjuster, the National Nosocomial Infections Surveillance (NNIS) System risk index,<sup>2</sup> achieves most of its separation through categorization of the risk associated with procedures, rather than with patients, who are classified as either "high

risk" (ie, an American Society of Anesthesiologists score of 3 or greater) or "normal risk." This dichotomous approach clearly adds value, although the categorization is often subjective and prone to misclassification.<sup>23</sup> The question that remains is, "Can we do better?" Clinical experience, a large literature identifying risk factors for infection, and surgeons with higher than average infection rates all tell us that we can. Russo and Spelman use their own experience in coronary artery bypass graft surgery to identify three patient-specific risk factors, obesity, diabetes, and noncardiac vascular disease, and then to create a risk index that predicts infection better than the NNIS System risk index. This study raises many important questions, including how much of the variation attributable to personal risk factors is captured by a risk index, how should one assess the generalizability of a new index, and how can one efficiently implement such an index when many institutions must obtain characteristics such as obesity by manual review of full-text medical records, if they are available at all?

Taken together, these three articles tell us much about what we know and what we don't know concerning the occurrence of SSIs. We certainly know that the answers to the questions "How many postoperative infections are there?" and "What is a patient's risk of infection?" depend on the definitions we use and where and how we search for them. We also know that we can substantially improve our ability to control for a patient's underlying risk of infection. And because of this, we can conclude that we will not be able to make meaningful comparisons—over time, between hospitals, and certainly between surgeons—until we can fairly characterize the riskiness of the patients as well as the procedures. Without such comparisons, we will have difficulty identifying best practices and we will not know the effect of our efforts to improve patient safety.

Therefore, we must continue to address the problems of infection detection and case-mix adjustment. However, we are likely to need additional approaches to these problems, because we are unlikely ever to have vastly greater resources, the prospect of much better health-care information systems (ones that know body mass index, for example) in the foreseeable future is uncertain, and postoperative care is likely to become even more dispersed.

To answer the most important questions, for the healthcare system as a whole, for hospitals, for surgeons, and for individual patients, we should develop additional benchmarks that can be uniformly applied to essentially all instances of a given procedure, without regard to a specific facility's infection control budget, the willingness of its surgeons to participate in a surveillance program, or the location of postoperative care. To accomplish this in the short-term, we will need to use data systems and information that already exist—hospital discharge data, the billing records that payors (insurance companies and health plans, including Medicare) routinely process, and outpatient antibiotic exposure data obtained from phar-

macy claims payment systems. These information systems have the advantage of being nearly universal, of using standard coding systems, and of being highly automated, so that the cost of extracting information about postoperative infections will be low. What will the trade-offs be for using such systems? We surely can't use these information systems to identify the same cases that traditional surveillance does, but there are suggestions that they can identify enough risk-adjusted outcomes of interest to serve as a meaningful performance measure for our healthcare system as a whole and for hospitals, if not for individual surgeons or patients.<sup>24-27</sup>

We will continue to need traditional methods for identifying SSIs, and we should strive to improve the accuracy and interpretability of these methods. In addition, we should consider nontraditional uses of the large amount of data that surround us, to obtain the information we need to make surgery as safe as possible.

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