Unnecessary Deaths: The Human and Financial Costs of Hospital Infections

3rd Edition

By Betsy McCaughey, Ph.D.





Essential Facts You *Need* to Know About Hospital Infections

Keep in mind:

- · We have the knowledge to prevent hospital infection deaths.
- · Yet most hospitals have failed to act.
- The situation is growing more dangerous because, increasingly, hospital infections cannot be cured with commonly-used antibiotics.

Essential facts:

- Infections contracted in hospitals are the fourth largest killer in America. Every year in this country, two million¹ patients contract infections in hospitals, and well over 100,000 die as a result, ² more deaths than from AIDS, breast cancer, and auto accidents combined.
- 2. A few hospitals in the U.S. too few are proving that infections are almost entirely preventable. How are they doing it? Through rigorous hand hygiene, meticulous cleaning of equipment and rooms in between patient use, testing incoming patients to identify those carrying dangerous bacteria, and taking precautions to prevent these bacteria from spreading to other patients.
- 3. Hospital infections add an estimated \$30.5 billion to the nation's hospital costs each year. Patients, insurers and taxpayers pay part of that, but hospitals have to absorb much of the cost. As a result, infections erode hospital profits. Preventing infections can turn a financially failing hospital profitable.
- 4. Until recently, infection was considered an inevitable risk of hospitalization. Now, there is compelling evidence that nearly all hospital infections are preventable when doctors and staff adhere to researchbased guidelines. This puts physicians, hospitals, and hospital board members in a new legal situation. The belief that

- infections are unavoidable shielded hospitals and doctors from liability for decades. But not in future. Hospital infections could be the next asbestos.
- 5. Better infection prevention in hospitals is essential to prepare the nation for avian flu or bioterrorism. If avian flu were to wing its way to the U.S., the death toll would depend largely on what American hospitals did when the first avian flu patients were admitted. If hospitals have effective infection controls in place, they can prevent bird flu from infecting other patients who did not come in with it. If not, bird flu could sweep through hospitals. Right now, most hospitals are woefully under prepared. Hospitals have failed to stop the spread of ordinary infections spread by touch and would not be able to contain flu viruses, which are communicated by droplets from coughing and sneezing as well as touch. Shoddy infection control is poor preparation for a flu epidemic and poor homeland security as well.
- 6. Hospital infection is a far deadlier problem than the number of uninsured. The Institute of Medicine estimates that as many as 18,000 people a year die prematurely because they don't have health insurance. That's tragic. But five times as many people die each year from hospital infections, and most of them are insured.³



Dear Reader,

This book has been written to help you and to enlist your help in correcting a deadly situation that kills well over 100,000 people in this country each year — more deaths than from AIDS, breast cancer, and auto accidents combined. Where does it kill? In our hospitals. What is it? Hospital infection.

The death toll is staggering. So is the economic cost. Hospital infections add over \$30 billion a year to what the nation spends on hospital care, enough to pay for a large part of the Medicare Part D drug program.⁴

These infections are almost all preventable. An increasing number of hospitals in the U.S. are proving it, reducing some of the deadliest types of infections by 90 percent. Their achievements prove that we have the knowledge to solve this problem. What is lacking is leadership.

That is why I founded the Committee to Reduce Infection Deaths (RID): to motivate hospitals to make infection prevention a top priority; to inform patients about the steps they can take to reduce their risk of infection; and to ensure that no matter where you live, you can find out which hospitals in your area have the worst infection problems.

- RID holds forums for hospital administrators, public health officials, law-makers, medical educators, insurers, and patient advocates, showing them how infections can be eradicated and how much money can be saved. The humanistic reasons to stop hospital infections are obvious. RID forums also make a compelling economic case for infection prevention, showing that for some hospitals, preventing infection can actually make the difference between profitability and loss.
- RID educates the public through television, radio, popular publications, and our website. One of our most important educational tools is the "15 Steps You Can Take to Reduce Your Risk of a Hospital Infection," which is included in this study.
- RID works with state lawmakers and other policy makers to develop hospital infection report cards, because if you need to be hospitalized, you should be able to choose a hospital with a low infection rate.
- RID provides the latest, most accurate research in a format that doctors, nurses and other caregivers can access electronically at any time, from any where. RID's electronically delivered curriculum bridges the gap between today's research and yesterday's practices.

■ RID is encouraging medical schools and nursing schools to educate their students about how bacteria are spread from patient to patient in hospitals and the precautions that should be taken to protect their patients — a subject that is almost entirely neglected in most schools.

RID has a distinguished advisory committee, including: Dr. Carlene Muto, Director of Infection Control and Hospital Epidemiology, University of Pittsburgh-Presbyterian Medical Center; Dr. Richard Shannon, Professor of Medicine, Hospital of the University of Pennsylvania; Dr. Jeffrey Borer, Chairman of the Division of Cardiovascular Pathophysiology at New York-Presbyterian Medical Center; Dr. Steve O'Brien, Vice Chairman of the Sports Medicine Department at Hospital for Special Surgery; Dr. Allen Hyman, Former Chief of Staff and Medical Director of New York-Presbyterian Medical Center; Dr. Bart Pasternack, a cardiovascular surgeon at Norwalk Hospital and Yale-New Haven Hospital in Connecticut; Dr. Alan Jasper, Chairman of the Department of Critical Care Medicine and Former Chief of Staff at St. Vincent's Medical Center in Los Angeles; Jane Barnsteiner RN, PhD, FAAN, Professor of Pediatric Nursing at UPENN School of Nursing; Dr. Sherwin Nuland, author of *A Doctor's Plague*; and Dr. Elizabeth Whelan, founder of the American Council on Science and Health. RID mourns the loss of our advisory board member Nobel Laureate Dr. Joshua Lederberg, who died on February 2nd, 2008. Other members include philanthropists and civic and corporate leaders.

Everyday you hear about health care problems such as providing for the uninsured. The Institute of Medicine estimated that as many as 18,000 people a year may die prematurely because they don't have health insurance. But consider this even more tragic fact: five times that many people die each year from hospital infection, and most of them are insured. Having insurance is no guarantee that you will be safe in the hospital. The only way to ensure that is to clean up this deadly problem.

Betsy McCaughey, Ph.D.

Founder and Chairman

Committee to Reduce Infection Deaths

www.hospitalinfection.org

Table of Contents

I.	Third World Hygiene in Our First Class Medical System			
II.	The Major Problem: Poor Hygiene4			
III.	MRSA Screening Is Essential			
IV.	Success Stories: Infections Can Be Eradicated			
	A. Dr. Carlene Muto Describes Victory over MRSA at the U. of Pittsburgh Medical Center 11			
	B. Dr. Barry Farr Recalls Early Victories at the University Of Virginia Hospital			
	C. Dr. Richard Shannon Aims for Zero Central Line Infections			
V.	Protecting Patients From Central Line Infections			
VI.	To Prevent C.diff, Cleaning Is Essential			
VII.	Preventing Infections Makes Hospital More Profitable21			
VIII	. Hospital Infection is the Next Asbestos			
IX.	Shouldn't Medical Students Be Taught Hygiene?			
	15 Steps You Can Take to Reduce Your Risk of a Hospital Infection			
X.	The Importance of Hospital Infection Report Cards			
App	endix A: RID's Model Hospital Infection Report Card Bill			
App	endix B: Model Screening Bill			
for P	endix C: Society for Healthcare Epidemiologists of America Guideline reventing Nosocomial Transmission of Multidrug-Resistant Strains of hylococcus Aureus and Entrococcus			
•	endix D: The Institute for Healthcare Improvement's 5 Million Lives Campaign			
	ut the Author54			
End	notes			

"We believe that healthcare facilities in the United States can be as successful in controlling MRSA as health care facilities in Northern Europe and Western Australia...."

– William Jarvis, December 2007 ⁶

Third World Hygiene in Our First Class Medical System

Every day in hospitals across the United States wondrous medical procedures rescue patients from the brink of death. But there's a catch: in these same hospitals, hygiene and procedures are so inadequate that up to 10% of hospital patients contract infections.⁷

Infections that have been nearly eradicated in some countries are raging through American hospitals. In 2003, the Society for Healthcare Epidemiologists of America warned that although hospitals have infection prevention programs, "there is little evidence of control in most facilities."

The danger is growing because hospital infections, increasingly, cannot be tamed with commonly-used antibiotics. One of the deadliest germs is methicillin-resistant Staphylococcus aureus (or MRSA). Patients who do survive MRSA often spend months in the hospital and endure repeated surgeries to cut out infected tissue. In 1974, 2 percent of Staph infections were MRSA. By 1995, the number had climbed to 22%, in 2003 an alarming 64%, and now even higher. In 2007, a research team led by William Jarvis concluded that MRSA is "has become

endemic in virtually all U.S. health care facilities."¹⁰

Denmark, Holland, and Finland once faced similar rates, but brought them down below 1 percent.11 How? Through rigorous hand hygiene, meticulous cleaning of equipment and rooms in between patient use, testing incoming patients for MRSA and other drug resistant bacteria, and taking precautions to prevent transmission to other patients. Wheelchairs and other equipment used to transport patients who test positive for MRSA are not used for other patients, and hospital staff have to change their uniforms and footwear after entering the rooms of MRSA patients, before they are permitted in other areas of the hospital.

An increasing number of hospitals in the United States are proving these precautions work here too. The University of Virginia Hospital eradicated MRSA.¹² The Veterans Hospital in Pittsburgh reduced MRSA by 85 percent in a pilot program.¹³ The University of Pittsburgh-Presbyterian Medical Center slashed MRSA by 90 percent in the medical intensive care units in a pilot program,¹⁴ and a

Yale-affiliated hospital in New Haven, Connecticut, cut MRSA infections by two thirds in a surgical intensive care unit. Beth Israel Medical Center in New York City reduced MRSA infections 65% from 2003 to 2007. Brigham and Women's Hospital in Boston reduced MRSA bacteremia 77% in intensive care and 67% hospital-wide. 16

Smaller hospitals around the nation are providing leadership as well. In November, 2007 Pitt County Memorial Hospital in South Carolina announced that screening all incoming patients and all elective surgery patients for MRSA, and then taking precautions to prevent the germ from spreading, achieved a 67% reduction in ventilator-associated pneumonia and a 60% drop in MRSA urinary tract infections in eight months. Pitt County is now hosting forums to show other hospitals in the region how it's done.

Twenty-nine healthcare institutions in Iowa eliminated another drug-resistant germ, *vancomycin-resistant Enterococcus* (or VRE).¹⁷ Unfortunately, most U.S. hospitals have not implemented these precautions.

How Many Hospital Infections? Tallying the Human Cost

Dad's death certificate said "organ failure," my friend recalled. "That's like saying the cause of death was dying. It was the hospital infection that killed him."

Until recently, the Centers for Disease Control and Prevention estimated that one out of every twenty patients (5%) contracts an infection in the hospital. That figure was based on what a small number of hospitals voluntarily report to the CDC's National Nosocomial Infection Survey (NNIS). But two new studies show that the CDC estimate understates the problem by as much as half.

First, a nationwide survey announced in June, 2007 found that 2.4% of inpatients were infected with hospital-acquired MRSA. That's just one bacterium! Surely the overall infection rate would be far higher than 5% if all bacteria were counted. The new survey was far more extensive that previous CDC collection efforts, and included 1,200 hospitals (nearly a quarter of the hospitals in the U.S.). Also, the CDC customarily relies on data from the ICU only, and the new survey found infections are more prevalent in other parts of the hospital than in ICUs.²⁸

Second, a report in the October 17, 2007 issue of the Journal of the American Medical Association made headlines by disclosing that more people in the U.S. die from MRSA than from HIV. The MRSA death toll is significantly larger than previous CDC estimates. Why? The new study examined patients' actual laboratory results, rather than basing its findings on what hospitals tell families, put on death certificates, or report to the CDC. In other words, the new report exposed the truth gap between actual cases and those reported.²⁹

If the same methodological improvements that result in far higher MRSA figures were used to tally hospital infections from all types of bacteria, instead of this one superbug, total infection rates would be found to be much higher than 5% and the annual death toll would be found to be substantially greater than 100,000.

\prod

The Major Problem: Poor Hygiene

 Λ stoundingly, physicians and other Π caregivers break the most fundamental rule of hygiene over half the time by failing to clean their hands before treating patients.¹⁸ Programs to encourage better compliance have been disappointing. Brigham and Women's Hospital in Boston assessed the impact of installing dispensers for alcohol based hand cleaners in every patient's room and conveniently in the hallways, and conducting a year long campaign on hand hygiene. The results? Hand cleaning temporarily improved from 40% to 80%, but quickly dropped back to 60%. 19

Unfortunately, caregivers often think putting on gloves — without cleaning their hands first — is sufficient. But pulling on gloves with unclean hands simply contaminates the gloves.

Cleaning hands is essential, but it's only the first step. Caregivers also need to learn how to prevent their hands or gloves from becoming re-contaminated before touching the patient. Stand in the emergency room, and watch caregivers clean their hands, put on gloves, and then reach up and pull open the privacy curtain to see the next patient. That curtain is seldom changed, and it is frequently full of bacteria. The result? Caregivers' gloves are soiled again.

As long as surfaces in hospitals are inadequately cleaned, caregivers' hands will become recontaminated seconds after they are washed.

How dirty are hospitals? Boston University researchers examining 49 operating rooms in four New England teaching hospitals found that more than half the objects that should have been disinfected were overlooked by cleaning staff. A follow-up study of patients' rooms in 20 hospitals in Washington, D.C., Connecticut and Massachusetts found that more than half the surfaces that are supposed to be cleaned after a patient is discharged and before the next patient is admitted were overlooked.²⁰

Research shows that nearly three quarters of patients' room are contaminated with MRSA and VRE.²¹ These bacteria are on cabinets, counter tops, bedrails, bedside tables, and other surfaces. Once patients and caregivers touch these surfaces, their hands become vectors for disease. One study showed that when a nurse walks into a room occupied

by a patient with MRSA and has no patient contact, but touches objects in the room, the nurse's gloves are contaminated 42% of the time when leaving the room.²²

MRSA and VRE can live for many hours on surfaces and fabrics. Thorough cleaning is highly effective in reducing the spread of drug-resistant bacteria to patients.

In a 2006 study, researchers at Rush University Medical Center in Chicago trained the environmental services staff to soak surfaces with detergent and wait, rather than merely spraying and quickly wiping, and to clean commonly overlooked objects such as telephones, remote controls and faucets. The result was a two-thirds reduction in the spread of VRE to patients.²³ Even the cash-strapped British National Health Service recognizes that intensive cleaning is a bargain compared with the cost of treating infections. By nearly doubling cleaning staff hours on one ward, and with no other interventions, a hospital in Dorchester, England reduced the spread of MRSA by 90%. The financial results were also impressive. Savings from infections avoided were 3.5 times the cost of the added cleaning.

Environmental surfaces are vectors for drug-resistant bacteria, but the most important sources of these bacteria are the patients coming into the hospital. Amazingly, most hospitals in the U.S. don't test incoming patients for MRSA. Seventy to ninety percent of patients carrying MRSA are unknown. They are the silent reservoir in the hospital. Knowing which

patients are sources of bacteria is the key to stopping the spread.²⁴

Recent research highlights the danger of MRSA and other superbugs lingering on surfaces long after the patient who carried the germ has been discharged. In one nine-bed ICU, more than half the patients who picked up MRSA after entering the ICU acquired a strain of the bacteria not present on other patients in the ICU at the time. In other words, the bacteria had been left behind on floors, bedrails, tables, and other surfaces by patients already discharged. These findings demonstrate 1) the importance of housekeeping and 2) how essential it is to know which patients entering the ICU are carrying the bacteria.

Placing a patient in a room or even a wheelchair previously used by someone who unknowingly carried MRSA put that patient at risk.

When hospitals fail to identify which patients are carrying superbugs, hospital uniforms and equipment become conveyor belts for infection. When doctors and nurses lean over a patient with MRSA bacteria, their white coats and uniforms pick up that bacteria 65% of the time, allowing it to be carried to other patients.²⁵

A 2007 University of Maryland study revealed just how contaminated the clothing worn by hospital personnel can be. In the study, sixty-five percent of doctors and other medical personnel treating patients admitted that they change their lab coat less than once a week, even though they know it is contaminated. Fifteen

percent admitted they change it less than once a month.²⁶

Hospitals that are conquering infections require their staff to put on fresh gowns or disposable aprons every time they approach the bedside of patients carrying MRSA, not just infected patients, but all patients carrying the bacterium. (The disposable aprons cost a nickel and are ripped off rolls like clear, plastic dry cleaning bags.)

Stethoscopes, blood pressure cuffs, pulse oximeters, wheelchairs, and other equipment are frequently carrying live bacteria. Do doctors clean the stethoscope before listening to a patient's chest? Not usually, though the American Medical Association recommends it.

When a nurse wraps a blood pressure cuff around the patient's bare arm, the cuff frequently carries live bacteria, including MRSA. In a September 2006 study, 77% of blood pressure cuffs rolled from room to room in the hospital were contaminated.

An astounding example of the impact of contaminated equipment occurred in a burn unit at an academic hospital in Galveston Texas. When the VRE outbreak began, hospital personnel cultured every surface in the unit. The results were revealing: 19% of bedrails, 26% of infusion pumps, and virtually all other surfaces were contaminated with VRE. Hospital personnel launched a "very aggressive" twice-daily cleaning of patients' rooms and equipment, including equipment that ordinarily was the responsibility of the nursing staff. Incoming patients

were cultured for VRE, and all caregivers wore gowns and gloves when treating positive patients. The "clean and screen" strategy succeeded. VRE was eradicated from the unit by October, 2000.

Then, in November, a physician treating a patient with a large shoulder and neck burn noticed one item had not been cultured — the EKG wire just used. Four days later, his patient's wound tested positive for VRE — and molecular typing confirmed the germ had come from the wire. Amazingly, the VRE on that wire had been left behind by a patient discharged 38 days earlier!

We have the knowledge to prevent infection. What has been

VRE Contaminates Surfaces in an Academic Hospital, Galveston, TX						
Site	(%)	Positive	for VRE			
Infusion pur	nps		26 %			
Bed rails			19 %			
Shelves			19 %			
Overbed tab	les		18 %			
Bedside tabl	es		17 %			
Pulse oxime	ters		14 %			
Stethoscope	S		13 %			
Monitors			13 %			
Suction cani	sters		12 %			
Chairs			11 %			
Doors			11 %			
IV poles			10 %			
Oxygen flov	mete	rs	8 %			
Faucets			6 %			
Miscellaneou	18		14 %			
Total			13.5 %			

BP Cuffs As Vectors of Disease

In 2003, a university hospital in Tours, France, examined 203 BP cuffs used in medical, surgical, ICU, and emergency units.

Type of BP Cuff	Total Number	% Contaminated
On Nurses' Trolleys	35	77%
Individual	41	63%
Wall Model	57	53%
Stored	52	17%
Newly Cleaned (with disinfecting detergent)	18	0%

Extensive contamination of BP cuffs (30% of contaminated cuffs carried MRSA).

Source: C de Gialluly et al., "Blood Pressure Cuff as a Potential Vector of Pathogenic Microorganisms: a prospective study in a teaching hospital," *Infection Control and Hospital Epidemiology* 27.9 (2006): 940-3.

lacking is the will. In 2003, a committee of the Society for Healthcare Epidemiologists of America codified the precautions that have worked well in Denmark, Holland, and Finland and in the hospitals here in the U.S. that have tried them. The essence of these guidelines (see Appendix C) is cleaning and screening. This twofold strategy works. One study shows that MRSA bacteria spread from patient to patient 16 times as fast under standard CDC precautions as under the more rigorous precautions outlined in the 2003 guidelines. What a shame that more hospitals are not implementing these lifesaving precautions. The next chapter will review the compelling evidence for screening.27



MRSA Screening Is Essential

What kills more than five times as many Americans as AIDS each year? Hospital infections. Yet federal officials at the Centers for Disease Control and Prevention, who are calling for voluntary blood testing of all patients to stem the spread of AIDS, are not recommending a test that is essential to stop the rapid spread of another killer sweeping through our nation's hospitals: MRSA.

On September 19, 2006 the Centers for Disease Control and Prevention recommended universal testing for HIV. One month later, a Centers for Disease Control and Prevention committee issued new guidelines to prevent hospital infections but chose not to recommend that hospitals begin screening all patients for MRSA.³⁰

Is the MRSA test more invasive than the HIV test? No, it's less invasive — a simple skin or nasal swab to determine which patients carry the bacteria.

Is the MRSA test more expensive? No. The rapid MRSA test costs about the same as the rapid HIV test, \$20 or so.

Is MRSA testing needed? Yes, because MRSA is transmitted easily

from patient to patient on clothing, medical equipment, hands, and gloves.

Research shows that you cannot prevent MRSA infections until you identify which patients bring these bacteria into the hospital. Patients who unknowingly carry MRSA shed it in tiny particles on bedrails, wheelchairs, blood pressure cuffs, stethoscopes, and the floor under their beds. They don't realize they have it, because the germ doesn't make you sick (infected) unless it gets inside your body via a catheter, a surgical incision or other open wound, or a ventilator.

Among developed nations, Japan and the U.S. have the worst records of failing to control the rapid rise of drug-resistant hospital infections.²⁹ Data from the Centers for Disease Control and Prevention indicate that MRSA hospital infections increased 32 fold from 1976 to 2003.³¹

For a decade, the Centers for Disease Control and Prevention have rebuffed calls for screening. In 1996, in the *Journal of the American Medical Association*, a panel of experts warned that hospitals faced an "unprecedented crisis"

due to drug-resistant infections.³² In 2003, a committee of the Society for Healthcare Epidemiologists of America warned again that screening patients at risk for carrying MRSA was urgently needed.³³

In 2004, Dr. John Boyce announced that screening had reduced MRSA infections by two thirds in an intensive care unit at a Yale-affiliated Connecticut hospital. Based on this study and others, Boyce and co-researchers concluded that patients will not be protected from MRSA until hospitals start screening.³⁴

That is the compelling conclusion of a 9 year study done at the Brigham and Women's Hospital in Boston and published in the fall of 2006 in *Clinical Infectious Diseases*.³⁵ Researchers found that installing dispensers of alcohol-based hand cleaners

in each patient's room and outside each patient's room had no significant impact on MRSA bacteremia rates. Similarly, a subsequent year-long hand hygiene education campaign achieved no statistically significant reduction in MRSA bacteremia. But initiating routine surveillance cultures for all ICU patients and contact precautions for patients testing positive for MRSA resulted in an impressive 75% drop in MRSA bacteremia in intensive care units and a 67% drop hospital wide.

Researchers called the infection reduction at Brigham and Women's "profound." They explained that the reduction could have been even greater had they either used a rapid MRSA test, instead of a culture that took two days, or preemptively isolated patients until their cultures came back from the laboratory. The two day

New British Recommendations

- 1. Screen all patients admitted to "high risk" units, such as the ICU, cardiothoracic, orthopedic, and burn units.
- 2. Minimize movement of MRSA-positive patients.
- 3. Use gowns and disposable aprons when treating MRSA-positive patients.
- 4. Launder privacy curtains or use disposable curtains.
- 5. Decontaminate trolleys and wheelchairs after patient use.
- 6. Before surgery, attempt to decolonize MRSA positive patients.
- 7. In the recovery area, segregate MRSA positive patients.

Source: Specialist Advisory Committee on Antimicrobial Resistance (established to advise the UK government), "Guidelines for the control and prevention of methicillin-resistant Staphylococcus aureus (MRSA) in healthcare facilities," *Journal of Hospital Infection* 635 (April 2006).

delay permitted some spread of bacteria from patients who unknowingly carried MRSA to other patients who did not come in with it.

Despite these and many other studies, the CDC continue to equivocate, rather than urging all hospitals to screen incoming patients for MRSA and take contact precautions to prevent the bacteria from spreading to other patients. "There are at least fifty studies demonstrating the effectiveness of these precautions," explains Dr. Carlene Muto, and "not one study suggesting it's possible to control MRSA without them."

Fortunately, some hospitals are leading the way, including Evanston Northwestern, the seventeen Veterans Affairs medical centers, New England Baptist Hospital in Boston, Johns Hopkins in Baltimore, North Shore University Hospital on Long Island, and Beth Israel Medical Center in New York City.³⁷ Even the cash-strapped British National Health Service has launched MRSA testing. Hospitals that don't screen are putting their patients at greater risk of an MRSA infection.

Now that the Centers for Disease Control and Prevention are telling hospitals to test for HIV, they should call for MRSA screening as well. Lax guidelines encourage hospitals to do too little. Every year of delay is costing thousands of lives and billions of dollars.

IV

Success Stories: Infections Can Be Eradicated

A. Dr. Carlene Muto Describes Victory Over MSRA at the University of Pittsburgh-Presbyterian Medical Center³⁸

66 Tt's a fabulous feeling," says Dr. Carlene Muto, reflecting on the team effort that has resulted in a 90 percent reduction in methicillin-resistant Staphylococcus aureus (MRSA) in the medical intensive care unit at the hospital where she is director of infection control. How long did it take? Three years. Ask her how it was done. She explains that it required total commitment from the top leadership at the hospital and caregivers. Many hospital administrators worry that they can't afford to implement these precautions. The truth is, they can't afford not to. Infections erode hospital profits, because rarely are hospitals paid fully for the added weeks or months of care when a patient gets an infection.

When Muto came to UPMC-Presbyterian, the flagship hospital in the University of Pittsburgh system, in the 1990s, drug-resistant *Staphylococcus aureus* was a rapidly growing problem. In 2000, Muto launched a campaign to

eradicate the "superbug" in the hospital's medical intensive care unit. Critical to the strategy was active surveillance culturing — meaning that every patient coming into the intensive care unit who might be carrying MRSA was cultured. Muto, one of the co-authors of The Society for Healthcare Epidemiologists of America's guideline, emphasizes that you can't eliminate infection until you know which patients are the sources of the bacteria. Every patient who tested positive was isolated, and doctors and nurses treating them wore gowns and masks, and kept equipment used on these patients away from others. By 2003, MRSA was almost eliminated. The strategy has worked so well that it has now been expanded to all 15 intensive care units in the hospital system.

The key, explains Muto, was to identify every patient carrying the dangerous bacteria. "We had total compliance, 98 percent to 100 percent, with culturing patients," she said, adding that she was astonished.

After all, asking nurses to culture every new patient in the ICU meant adding one more thing to an already long list of tasks they have to do. The staff reaction, says Muto, "has been overwhelmingly positive." "That's essential," she adds. "You can come up with an idea, but no matter how great it is, you have to have the buy in from the staff at the point of care."

Getting caregivers to clean their hands has been a tougher challenge, in part because at the beginning, Muto explains, some "nurses didn't realize that if they went into a room of a patient in isolation and didn't touch the patient or the bed linen but did touch other surfaces such as countertops, their hands were contaminated."

Now that the education process is well under way, hand cleaning

compliance is about 69 percent, well above the national average but not good enough for Muto and her team. The top leadership at UPMC-Presbyterian is taking an uncompromising position on the failure of staff and doctors to clean their hands. The hospital is getting set to impose stiff penalties, including firing staff members who chronically ignore hand cleaning rules and denying doctors the privilege of practicing at the hospital.

The goal? "Our goal is 100 percent compliance with hand cleaning, 100 percent compliance with gowning, 100 percent compliance with surveillance culturing," says Muto, adding excitedly that she can only imagine what can be achieved when they reach perfection.

B. Dr. Barry Farr Recalls Early Victories³⁹

Barry Farr remembers the first outbreak of MRSA at the University of Virginia Hospital. It was 1978, and Farr and his wife had recently come to the hospital to train, having just finished medical school. "MRSA was wildly out of control," he recalls, and the hospital was doing "what most American healthcare facilities are still doing today." As a result, the hospital "failed miserably to control the MRSA."

For nearly three years, as the outbreak raged on, the hospital followed a business-as-usual approach: no routine cultures were being taken to identify the patients silently carrying

the bacteria. The result, recalls Farr, was that doctors were touching patients who had MRSA, or allowing their white coats to brush up against them, and then passing the bacteria on to other patients without knowing it. At the hospital infection control meetings, the mood was pessimistic and apathetic. Staff members were saying "no one has ever controlled this."

Finally, after three years of failure, the hospital took a radical step, inspired by the success of several European countries that had brought MRSA under control. The hospital began regularly testing patients for

the bacteria and isolating those who tested positive. The results were stunning. Soon after the testing began, in December 1980, MRSA declined rapidly, and by the summer of 1982, the hospital was MRSA free. "It was beautiful," Farr recalls.

Surveillance culturing — identifying every patient carrying the bacteria — was the key to thwarting the outbreak and eradicating MRSA, says Farr. It was to work again a decade later.

The University of Virginia hospital was struck with MRSA a second time in the early 1990s, when a surgeon apparently walked into the neonatal intensive care unit with MRSA on his hands or clothing and transmitted it to one of the babies. Quickly it was spread to babies in the neighboring bassinettes, and then to another neonatal intensive care unit when one of the babies carrying the bacteria was moved there. The hospital immediately put into place the same precautions that had worked a decade earlier, and the outbreak was curtailed. Culturing every baby, and isolating every one who tested positive, was once again the key.

Would this method conquer other deadly bacteria as well? Soon afterward, the hospital faced an outbreak of *vancomycin-resistant Enterococi* (VRE), which spread rapidly to 30 percent of patients on eight separate wards. After several months, the hospital brought the outbreak under control once again by testing patients, isolating the carriers, and making sure that all staff put on gowns and gloves when treating them.

Are the University of Virginia's successes atypical? "No," says Farr. "There are over ninety studies, probably 100 by now," demonstrating that this method works. Yet antibiotic-resistant infections are "clearly out of control in the American health care setting." Why? Farr suggests that faulty cost-cutting is partly to blame.

Hospital administrators complain about the cost of these rigorous precautions, but the data proves these precautions save money. Farr compared the University of Virginia hospital with several other university hospitals of similar size. These other hospitals "are spending between \$1 million and \$3 million a year extra to treat antibiotic-resistant infections. far more than what UVA has had to spend on gowns, cultures, and gloves. We're taking the ounce of prevention approach. Many other hospitals are taking the pound of cure approach."

Another reason few hospitals are adopting rigorous infection control is that the public has not demanded it. "In Britain there is a public outcry over the failure to control MRSA infections in hospitals, and the British government is reportedly now considering firing hospital directors that fail to take effective measures to control MRSA," says Farr. "In this country there has been comparatively little outcry from the public and no urgent demands from the government that the spread of infections be stopped."

C. Dr. Richard Shannon Aims for Zero Central Line Infections

When Dr. Richard Shannon told the top executives at Allegheny Hospital that he wanted to do something about central line-associated blood stream infections (CLABs), the hospital leadership expected him to suggest reducing them by 10 or 20 percent over several years. To their surprise, Shannon said he wanted to totally eradicate these deadly infections in ninety days. And he did it! Even more amazing, he and his staff kept these infections near zero in the medical intensive care unit and coronary care unit during the entire next year, achieving a 95 percent reduction in CLAB-related deaths.40

Why strive for merely minor improvement when lives are at stake? Shannon's pet peeve is benchmarking — the thinking all too common in hospitals today that it's okay to have infections and medical errors so long as they don't exceed the national average. "Who volunteers to have a family member get one of the infections we plan on having this year?" The goal has to be zero infections and perfect care, says Shannon, who is Chairman of the Department of Medicine at Allegheny. ⁴¹

How was that goal reached? By ensuring that all caregivers meticulously follow a regimen for inserting and removing central lines that includes masks, gowns, gloves, and drapes; inserting lines in the neck area rather than in the groin

area, which is more difficult to keep clean; rearranging supply closets to ensure that the supplies needed to carry out this regimen are easily accessible, even when staff are rushed; and empowering all staff members to enforce hand cleaning and other rules of hygiene. If a doctor doesn't clean his hands, the nurse working alongside can call a halt to the procedure until the doctor complies.

Shannon oversees some employees and a \$150 million budget. Nevertheless, he makes time to speak across the country, with PowerPoint in tow, showing his audiences that preventing infections is possible and profitable. Doing the right thing costs less, he says, using Allegheny's financial records to prove the point. A typical example is the tragic case of a woman who came into the hospital for stomach reduction surgery, a procedure that should have produced a \$9,900 gross profit for the hospital. But when she developed a central line-associated bloodstream infection and had to spend 47 days in the hospital, that profit turned into a \$16,000 loss. Preventing CLABs saved Allegheny \$1.4 million the first year.42

The best news of all is that the success at Allegheny is being duplicated by at least a few other institutions. At Johns Hopkins, catheter-related bloodstream infections in the

intensive care unit have been virtually eliminated. How? ICU staff were educated about the seriousness of catheter-related infections: a catheter-insertion cart was created to ensure that necessary equipment was readily at hand; doctors were asked daily whether catheters should be removed: bedside nurses were given a safety checklist to follow during insertion; and nurses were empowered to stop procedures if safety rules were not being followed. Peter Provonost, the intensive care physician at Johns Hopkins who developed the safety checklist, sees the success as proof that infections are not inevitable.43

That is Richard Shannon's mantra as well. Shannon is amazed that so little is being done nationwide to curb bloodstream infections and to halt the alarming rise in MRSA. Shannon asks why the procedures that reduced Staph infections by 85 percent in a pilot program at the V.A. Hospital in Pittsburgh are not being duplicated everywhere. "What if you had a patient with TB or SARS? Wouldn't you pull out all the stops, gloving and gowning and washing up all the time? Well, we haven't seen TB in years, and we've never seen SARS, but we have MRSA silently stalking us every day." The magnitude of the problem, he says, is "a call to action for all health-care providers to step up and get serious about all hospitalacquired infections."44

V

Protecting Patients from Central Line Infections

One of the deadliest hospital infections is the central-line associated blood stream infection (CLABS). At least 12% of patients who contract these infections die from them, and in some hospitals the mortality rate is above 40%. Astoundingly, new research shows that these infections are preventable. Here is the proof—at individual hospitals, in a large metropolitan area, and even across the entire state of Michigan.

CLAB Prevention Honor Role

■ Johns Hopkins

Baltimore, Maryland

In one ICU, Dr. Peter Pronovost and Dr. Sean Berenholz pioneered the interventions that eradicated CLABs. The results: A zero rate of central line infections and an estimated 43 infections and 8 deaths prevented from 1998–2002.⁴⁶

■ Michigan

103 ICUs in Michigan (85% of all ICU beds in the state), including academic hospitals and small community hospitals duplicated the Johns Hopkins method. The results? An impressive 66% reduction in CLABs with sustained

improvement over 18 months.⁴⁷

■ CLAB Quality Improvement Collaborative New York City

This regional effort involving 56 ICUs at 38 hospitals was launched in June, 2005. Early results indicate a 58% reduction in CLABs in ten months. At Beth Israel Medical Center, Dr. Brian Koll reports that both the critical care unit and the ICU have been CLAB free for almost two years!⁴⁸

How Are They Doing It?

- Educating caregivers about best practices when inserting lines.
- Ensuring caregivers use full barrier precautions (masks, gowns, gloves, and full sterile drapes) to insert lines.
- Inserting lines in the neck-shoulder area (subclavian) rather than in the groin area (femoral), which is more difficult to keep clean.
- Cleaning skin with chlorhexidine before insertion.
- Rearranging supply closets and pre-packaging supplies in kits to ensure that the regimen can be followed even when staff are rushed.

- Asking daily whether the line can be removed.
- Empowering caregivers to enforce sterile procedures and halt a procedure if they see a lapse.

Can Hospitals Afford To Prevent CLABS? They Can't Afford Not To!

- The statewide reduction in CLABS in Michigan was achieved with no additional ICU staffing.⁴⁹
- At Johns Hopkins, researchers estimated that the pilot program averted \$1.9 in treatment costs.⁵⁰
- "At Beth Israel Medical Center, says Dr. Brian Koll, total additional costs to launch the CLABS initiative were \$30,000 (to package the insertion supplies into kits); treatment costs avoided were \$1,500,000 not a bad investment!"⁵¹

A Below-Average Infection Rate Is Not Good Enough!

"We demonstrated that it is possible to nearly eliminate catheter-related blood stream infections; therefore we should not accept National Nosocomial Infection Surveillance mean values as a measure of success, but rather we should shift our focus to zero harm." — Sean Berenholtz, M.D., (*Critical Care Medicine*, 2004)⁵² Hopkins.

Treated Catheters Are an Important Supplemental Tool

Sterile procedures are the first line of defense against central line infections. But humans are fallible. New technology can help kill germs that manage to get into a central line. For patients who need a central line for five days or longer and are at high risk for infection, catheters treated with chlorhexidine and silver sulfadiazine or with the antibiotics rifampin and minocycline are recommended by the Agency for Healthcare Research and Quality. These devices complement rigorous attention to sterile procedures. Recent research at Stanford University shows that that although treated catheters cost twice as much as untreated catheters, they are cost-effective whenever they reduces infection risk by at least 30%, as numerous studies indicate they do.

VI

Preventing "C. diff": Cleaning Is Essential

Hygiene is the best defense against today's superbugs, MRSA and VRE. But it is also the best known shield against the next germ threat, Clostridium difficile or "C. diff." C. diff. killed more patients in England in 2006 than MRSA,⁵³ and the same hyper-virulent strain, dubbed ribotype 027, has invaded some hospitals in the U.S. and Canada. In fact, despite almost no news coverage until 2007, C. diff has been causing trouble for several years.

The Centers for Disease Control and Prevention tracked a nearly two-fold increase in C. diff infections from 1996 to 2003.⁵⁴ Two statewide studies in Oregon and Massachusetts found C. diff infections increasing at an even faster pace.⁵⁵ In the Montreal area of Canada, C. diff increased fivefold from 1997 to 2004.⁵⁶ Worse still, in both Canada and the U.S., the mortality rate from this disease is rising. Therefore, it is more important than ever to prevent it with rigorous hygiene, education of caregivers, and prudent use of antibiotics.

So what do we need to know about this bacterial villain? Outside of hospitals, it is normally found in the gastrointestinal tracts of about 5% of the general population. It doesn't usually cause trouble because other bacteria keep C. diff from getting out of control. In hospitals, the story changes. When a patient is put on antibiotics, the balance of bacteria in his gastro-intestinal system is affected, and C. diff. may take over, causing severe, watery diarrhea and inflammation of the colon.⁵⁷

It's the out of control nature of watery diarrhea that allows C. diff to spread so fast in a hospital. Although a small number of patients come into the hospital with C. diff spores in their bodies, many more ingest the germ through oral-fecal contamination, meaning traces of one patient's feces enter another patient's mouth. How could such a thing happen? The only answer is inadequate cleaning. Patients pick up the C. diff spores off contaminated bedrails, IV poles, tables, and other surfaces, virtually anywhere their hands can reach. Then they touch their lips, or touch their food and swallow C.diff along with their dinner roll. Caregivers unwittingly carry C. diff spores on their hands, uniforms, and equipment from patient to patient.

A 2006 study in the Journal of

Hospital Infection showed that onethird of blood pressure cuffs rolled from room to room carried C. diff spores on the inside of the cuff.⁵⁸ It's a short trip from a patient's arm to their fingertips and their mouth. Occasionally patients also get C. diff from inadequately cleaned rectal thermometers and endoscopes.⁵⁹

Environmental cleaning is so important that when it is not done regularly and rigorously, placing a patient in a room previously occupied by a patient with C. diff can be a fatal mistake. At Thomas Jefferson University Medical Center in Philadelphia, where C. diff was raging, three patients occupying the same room consecutively came down with C. diff. One died as a result.⁶⁰

In July and August, of 2005, eight infants in the neonatal intensive care unit at Intermountain Healthcare in Provo, Utah contracted C. diff. All eight infected infants had shared one of three beds in a corner of the NICU. The longer the hospital stay and the closer one is to a patient with C. diff, the greater the risk of contracting it.⁶¹

Training environmental services staff on how to clean more thoroughly is essential. At Case Western Reserve and the Cleveland VA Medical Center, researchers cultured commonly touched surfaces such as bed rails, telephones, call buttons, toilet seats, and bedside tables in the rooms of patients with C. Diff. After routine cleaning, 78% of the surfaces were still contaminated with C. diff spores. But once researchers disinfected the rooms with bleach,

including surfaces commonly overlooked by cleaners, only 1% of surfaces were still contaminated.

Dr. Carlene Muto and her colleagues at the University of Pittsburgh Medical Center - Presbyterian faced a 400% increase in C. diff infections in the year 2000. They responded with a comprehensive strategy that emphasized rigorous cleaning with bleach and rapid identification and isolation of C.diff positive patients to prevent the bacteria from spreading to other patients. (Additional interventions included reliance on soap and water rather than alcohol-based sanitizers to clean caregivers' hands, and controlled use of antibiotics beginning in 2003). This comprehensive strategy worked. By 2006, C. diff rates were down 71%, and severe cases of C. diff associated diarrhea fell by 89%.

At Intermountain Healthcare, after the eight infants contracted C. diff, the affected corner of the NICU was "cleaned from top to bottom," according to researchers there, including rockers and scales. "We launched extensive staff education related to C. difficile and its ability to be found on environmental surfaces," and "the importance of washing hands with soap and water when caring for a patient with C. difficile," they reported. The results? Not one new case of C. diff in the NICU in the next two years. 62

Educating hospital personnel on how patients are exposed to C. diff spores is essential. A study at one hospital found that resident physicians and other medical personnel were woefully under informed about

C. diff. For example, 39% didn't know that C. diff spores could be transmitted from patient to patient on equipment such as stethoscopes and blood pressure cuffs. Nearly 20% incorrectly thought C. diff was a blood borne pathogen, and almost 9% incorrectly believed it was transmitted through the air. Only about one third of medical professionals knew that cleaning hands with soap and water was essential, because alcohol sanitizers are often ineffective against C. diff. 63 This knowledge gap is dangerous to patients and costly to hospitals.

Based on an assessment of the increased length of stay required to treat C. diff patients in Massachusetts, researchers estimated "conservatively" that in 2005 alone, treating C. diff added \$3.2 billion to the cost of treating hospital patients nationwide.⁶⁴

Cleaning the hospital environment, educating personnel about C. diff., and controlling antibiotic use are essential to meet the C. diff challenge. In addition, hospitals need to consider other strategies. One is rigorous hand hygiene for patients. Nonambulatory patients are frequently handed a food tray, but have no way to clean their hands before dining. Their hands are contaminated with C. diff spores, which they ingest as they eat. Whenever and wherever C. diff threatens, patients need to be helped to clean their hands routinely before meals.

Looking Ahead

Though more research needs to be done, preliminary results suggest that adding a lactobacillus acidophilus milk product to the daily diet of patients on antibiotics may be effective at reducing antibiotic-associated diarrhea (AAD), including diarrhea caused by C. diff. A double blind study suggests that certain probiotics are effective, compared with a placebo, in reducing the incidence of antibiotic associated diarrhea by about half in patients on a variety of antibiotic regiments. The study suggests that some specific organisms may help restore the normal balance of bacteria in the gastro-intestinal system to spare patients from life-threatening diarrhea. If more research confirms these initial findings, hospitals may want to consider adding a nutritional supplement routinely to the diets of patients on antibiotics.

VII

Preventing Infections Makes Hospitals More Profitable

Many hospital administrators worry that they can't afford to implement these precautions. The truth is, they can't afford *not* to. Infections erode hospital profits, because hospitals are rarely paid fully for the added weeks or months of care when a patient gets an infection.

For example, Allegheny General

Hospital in Pittsburgh would have made a profit treating a 37-year-old video programmer and father of four who was admitted with acute pancreatitis, but the economics changed when the patient developed an MRSA bloodstream infection. He had to stay in the hospital 86 days, and the hospital lost \$41,813, according to research by Richard Shannon,

Estimated Hospital Costs of Hospital-Acquired Infection in the United States

2,000,000

Estimated infections per year

X

\$15,27569

(Average additional hospital costs when a patient contracts an infection)

= \$30.5 Billion

Per year spent treating hospital infections

Note: This figure does not include doctors' bills, home nursing bills, home nursing care, lost time at work, and other non-hospital costs.

former chairman of the Department of Medicine at Allegheny.⁶⁵

Similarly, a woman came into the hospital for stomach-reduction surgery, a procedure that should have produced a \$5,900 gross profit for the hospital. But when she developed a central line-associated bloodstream infection and had to spend 47 days in the hospital, that profit turned into a \$16,000 loss.⁶⁶

At Allegheny General Hospital, the average payment for a patient who developed a central line-associated bloodstream infection (CLAB) was \$68,894, but the actual average cost of treating the patient was \$91,733, leading to a gross loss of \$26,839 per case. The hospital had 54 such cases in the medical intensive care unit and coronary care unit between July 2002 and June 2005. The infections resulted in a total economic loss to the hospital of \$1,449,306.⁶⁷

Hospital infections add more than \$30 billion annually to the nation's health tab in hospital costs alone.⁶⁸ The tab will increase rapidly, as more infections become drug-resistant.⁷⁰

A new study based on all the hospital infections reported in Pennsylvania in 2005 dramatizes this enormous economic burden. The average charge for patients who developed an infection (\$173,206) was nearly four times as high as for patients admitted with the same diagnosis and severity of illness who did not contract an infection (\$44,367). The 11,688 infections reported added over two billion dollars in hospital charges that year. That's in one state alone!⁷¹

Other studies on the cost of infections found that:

- Post surgical wound infections more than double a patient's hospital costs. When a patient develops an infection after surgery, the cost of care increases 119 percent, on average, at a teaching hospital, and 101 percent at a community hospital.⁷²
- Urinary tract infections increase a patient's hospital costs by **47** percent at a teaching hospital and **35** percent at a community hospital.⁷³
- The average ventilator-associated pneumonia infection (a type of infection contracted when a patient is on a respirator) adds \$40,000 to a patient's hospital costs ⁷⁴
- Staphylococcus aureus infections are especially costly. According to a recent nationwide study, patients with Staph infections incur hospital costs that amount to more than triple the average hospital costs of other patients.⁷⁵

Not worried because your hospital's infection rate is well below the national average? Even hospitals with a below-average infection rate lose money on infections. A recent survey of 55 hospitals, where the infection rate averaged only 4.09% — well below the national average — showed that treating these infections wiped out inpatient operating profits.⁷⁶

The fact that hospitals lose money on infections doesn't necessarily prove that spending *more* on prevention will increase profits. Fortunately, there is compelling evidence that testing patients for drug-resistant bacteria and treating those who test positive with contact precautions yields a high return immediately and requires no capital outlays.

For example, Dr. Carlene Muto at the University of Pittsburgh-Presbyterian, where MRSA infections were slashed 90% in a pilot program, found that implementing these precautions in one medical intensive care unit cost \$35,000 in additional labor and materials, but prevented infections that would have cost over \$801,000 to treat. That's a 20 to 1 financial return the first year, not to mention lives saved.

Two community hospitals in Charleston, South Carolina, demonstrated that targeted surveillance — testing only patients deemed at high risk, such as patients recently hospitalized, living in a nursing home, or with kidney problems produces more modest reductions in infection and lower financial returns. This is not surprising, because a significant number of patients carrying MRSA go undetected. The costs of targeted surveillance, including laboratory tests and supplies such as gowns and gloves, cost \$113,955 and yielded just over a 10 to 1 return, saving the hospitals \$1,548,740 in avoided treatment costs.77

A recent review in Lancet concludes:

"Virtually all published analyses that have compared costs of screening of patients on admission and using contact precautions with colonised patients with the cost savings made by preventing health-care associated MRSA infections have concluded that the combination of surveillance cultures and barrier precautions results in cost savings for hospitals."

"THE COSTS
OF CARING
FOR PATIENTS
WHO BECOME
INFECTED WITH
MRSA ARE MUCH
GREATER THAN
THE COSTS OF
SCREENING
PROGRAMMES."78

A Model for Conducting Your Own Study

(From the U. of Pittsburgh-Presbyterian Medical Center)

Components of the Cost of Implementing Active Surveillance Culturing and Barrier Protections in One Medical Intensive Care Unit:

■ Laboratory: \$8,275

■ Personnel Time to Collect Samples: \$8,400

■ Supply Costs for Barrier Protections: \$16,337

■ Personnel Cost in Time to Don Protections: \$2,069

■ One Time Cost for Isolation Boxes: \$600

Results

Presumes 22% Annual Increase of Infections Without Intervention:

\$2,015,919

Cost of Expected HA-MRSA w/o Intervention

\$35,281

Cost of Interventions

=

\$1,980,638 SAVINGS

SAVINGS

Presumes a Stable Infection Rate Without Intervention:

\$801,652

Cost of Expected HA-MRSA w/o Intervention

\$35,281

Cost of Interventions

\$766,371

SAVINGS

Source: CA Muto et al., "Cost Avoidance Associated with Control of MRSA (University of Pittsburgh-Presbyterian Medical Center) — Presented at SHEA's 16th Annual Scientific Meeting (March 2006).

Staphlococcus aureus infections (both MSSA and MRSA) cost \$14.5 billion to treat in U.S. hospitals in 2003, according to a recent study by Gary Noskin and Robert J. Rubin. An estimated 85% of those infections were hospital acquired.⁷⁹

The federal Medicare program announced that beginning October, 2008, it will no longer pay the extra cost of treating several types of hospital infections, including device-associated urinary tract infections, central line infections, and certain types of post-cardiac surgical site infections. (The rule excludes all MRSA infections.) The new rule prohibits hospitals from billing patients or third parties for these costs. Federal officials indicated that the rule will probably be expanded to include other types of infections. Already several private insurers, including United Health Group and Cigna Corp. say they are exploring similar policies.80

This change in reimbursement should motivate hospitals to pursue the low-cost methods already available to prevent the most common infections, those caused by in-dwelling urinary tract catheters. Amazingly, most hospitals do almost nothing to prevent these infections. A survey of 719 hospitals across the nation showed that fewer than 25% monitored how long urinary tract catheters are left in, though monitoring is low-tech, almost cost free, and effective in reducing infection rates.⁸¹

VIII

Hospital Infection Is the Next Asbestos

Intil recently, infection was considered an inevitable risk of hospitalization. Now, there is compelling evidence that nearly all hospital infections are preventable when doctors and staff adhere to research-based guidelines. This puts physicians, hospitals, and hospital board members in a new legal situation. The belief that infections are unavoidable shielded hospitals and doctors from liability for decades. But not in future. Hospital infections could be the next asbestos.

Most victims who sue will not be able to prove precisely how the bacteria entered their body while they were hospitalized. Soon, it may not matter. Jurors will be told that the defendant physicians and hospital failed to implement guidelines provided by the federal Centers for Disease Control and Prevention (CDC) or such groups as the Society for Healthcare Epidemiology of America (SHEA), the Infectious Diseases Society of America (IDSA), and the Institute for Healthcare Improvement (IHI). Consequently, the argument will go, these defendants should be deemed negligent and held liable for patients' infections.

In 2006, the American College of Surgeons held a mock trial to anticipate the outcome of such a trial. In the mock trial, a hospital and physicians were sued when a patient suffered a catheter-related bloodstream infection and endocarditis that contributed to his death. The trial revealed that the resident physician who inserted the central line failed to wear a cap or gown, use a full sterile drape, or use an antimicrobialimpregnated or antiseptic-coated central line, even though the patient was at high risk for line sepsis and the line was expected to be in place for a prolonged period. Care did not meet CDC level 1A and 1B recommendations. The surgeons' defense attorney argued that the defendants were highly qualified physicians whose actions were within the standard of care in their community. It was also argued that only a small minority of hospitals and physicians follow CDC recommendations.

But mock trial experts warned that "community standard of care" is an inadequate defense if hospitals and physicians fail to implement recently published, nationally available, best practices. Physicians will have to read published guidelines and request that hospitals have the equipment on hand to implement them. The fact that the defendant hospital in the mock trial did not have appropriate central line kits in the intensive care unit was no excuse, trial experts agreed, because the surgeons had never requested a change of equipment. "Hospitals will not accept blame for stocking inappropriate equipment unless there is a clear paper trail that physician requests for proper supplies were denied." 82

In the real world, what must be done to avoid liability for infections? Ultimately, the courts will decide. Physicians and hospitals will need to make every effort to comply with published standards of infection prevention. I urge you to examine appendices C & D in the back of this book, as well as information provided by the Committee to Reduce Infection Deaths (www.hospitalinfection.org). For example, a hospital that fails to screen at least high-risk patients for MRSA colonization, despite a rapid rise in MRSA infections at the institution, may be at legal risk.

Even when a hospital complies with best practices, some plaintiffs' attorneys will still argue that a patient's infection is evidence enough that caregivers breached their duty. Every law student learns about the barrel that fell out of a merchant's second story window, injuring a customer below. In this textbook case, the merchant was held liable because the accident was itself evidence of

negligence, under the doctrine of res ipsa loquitor. Similarly trial lawyers will claim that an infection "speaks for itself" and shift the burden onto the hospitals and physicians to offer evidence that all best practices to prevent infection were in place.

Three recent events show how rapidly the legal environment is changing: In December 2004, Tenet Healthcare Corporation agreed to pay \$31 million to settle 106 individual lawsuits, filed between 1997 and 2002, on behalf of patients who suffered post-surgical infections after cardiac surgery at Palm Beach Gardens Medical Center.

Since then, thirteen lawsuits have been filed against Martin Memorial Hospital in Florida by patients who claim they contracted methicillinresistant Staphylococcus aureus (MRSA) infections while at the hospital for surgery. One is a wrongful death suit filed by a deceased patient's wife. In statements to the press, hospital officials claim that drug-resistant infections are a problem at all hospitals, and that their infection rate is below the national average.

That argument is also being used by Jewish Hospital in Louisville, Kentucky, which is facing numerous lawsuits. Among those are 69 filed by one lawyer, Joseph White, who has made unclean hospital conditions an attention-getting, graphic, and sympathetic explanation for why his clients contracted infections. The complaints include photos of unclean patient's bathrooms and testimonials of inadequately cleaned operating rooms with traces of dried blood on the

floor. Litigants even staged protests outside the hospital for evening news cameras, to claim that unsanitary conditions caused them to contract bacterial infections. The lesson is that physicians and hospitals will be put on the defensive even when facilities simply look unclean.

An attorney for Jewish Hospital told the press, "This is not going to become the asbestosis of hospital litigation." That may be wishful thinking.

The Committee to Reduce Infection Deaths would like to help hospitals protect their patients from infection and avoid lawsuits. Lawsuits are not the best way to improve patient care. They often result in unfair verdicts, and few truly injured patients ever make it to court (less than 2% according to the Harvard Medical Practice Study). Hospitals that act decisively to implement infection prevention guidelines will have the best insurance against costly damage awards.

IX

Shouldn't Medical Students Be Taught Hygiene?

That else needs to be done? Medical schools should be teaching future doctors the precautions they must take to protect their patients from infection. It's hard to believe, but most medical schools devote virtually no time, not even one full class, to showing students how bacteria are transmitted from patient to patient on clothing, equipment, and gloves, and what specifically they should be doing to prevent it. Dr. Frank Lowey, a professor at the New York-Presbyterian Hospital at the Columbia University Medical Center says, "it's something we should have done quite a while ago." Lowey says it's ironic that "there are curriculum committees devoted to making sure that bioterrorism is covered, and the risk of nosocomial infections far outweighs that."83

Some medical schools are stressing the importance of curbing the use of antibiotics.⁸⁴ That's good, because overuse of antibiotics wastes money and causes bacteria to morph into new, drug-resistant strains. But limiting the use of antibiotics won't stop hospital infections. Patients who

contract MRSA get it from unclean hands or contaminated equipment or clothing, not simply from taking antibiotics. No hospital has ever eradicated infection merely by controlling the use of these drugs.

When medical students put on their white coats and swear the Hippocratic Oath, they should be taught *how* to do no harm. Preventing the spread of bacteria is an essential part of that lesson. They should learn it before they go out on the hospital floors and touch their first patient.

15 Steps You Can Take to Reduce Your Risk of a Hospital Infection

- 1. Ask that hospital staff clean their hands before treating you, and ask visitors to clean their hands too. This is the single most important way to protect yourself in the hospital. If you're worried about being too aggressive, just remember your life could be at stake. All caregivers should clean their hands before treating you. Alcohol-based hand cleaners are more effective at removing most bacteria than soap and water.85 Do not hesitate to say the following to your doctor or caregiver: "Excuse me, but there's an alcohol dispenser right there. Would you mind using that before you touch me, so I can see it?" Don't be falsely assured by gloves. Gloves more often protect staff than patients. If caregivers have pulled on gloves without cleaning their hands first, the gloves are already contaminated before they touch you.86
- 2. Before your doctor uses a stethoscope to listen to your chest, ask that the diaphragm (or flat surface of the stethoscope) be wiped with alcohol. Numerous studies show that stethoscopes are often

- contaminated with *Staphylococcus aureus* and other dangerous bacteria, because caregivers seldom take the time to clean them in between patient use.⁸⁷ The American Medical Association recommends that stethoscopes routinely be cleaned for each patient. The same precautions should be taken for many other commonly used pieces of equipment too.
- 3. If you need a "central line" catheter, ask your doctor about the benefits of one that is antibiotic-impregnated or silver-chlorhexidine coated to reduce infection.⁸⁸
- **4.** If you need surgery, choose a surgeon with a low infection rate. Surgeons know their rate of infection for various procedures. Don't be afraid to ask.
- 5. Beginning three to five days before surgery, shower daily with 4% chlorhexidine soap. Drug stores that don't stock chlorhexidine soap are generally happy to order it for you. You don't need a prescription. One of the easiest brands to find is Hibiclens. Using this soap

will help remove any dangerous bacteria you may be carrying on your own skin that could enter your surgical incision and cause an infection.⁸⁹ Keep the soap away from your eyes and ears.

- **6.** Ask your surgeon to have you tested for *Staphylococcus aureus* at least one week before you come into the hospital. The test is simple, usually just a nasal swab. About one third of people carry Staphylococcus aureus on their skin, and if you are one of them, extra precautions can be taken to protect you from infection, to give you the correct antibiotic during surgery, and to prevent you from transmitting bacteria to other.⁹⁰
- 7. Stop smoking well in advance of your surgery. Patients who smoke are three times as likely to develop a surgical site infection as non-smokers, and have significantly slower recoveries and longer hospital stays.⁹¹
- **8.** On the day of your operation, remind your doctor that you may need an antibiotic one hour before the first incision. For many types of surgery, a pre-surgical antibiotic is the standard of care, but it is often overlooked by busy hospital staff.⁹²
- **9.** Ask your doctor about keeping you warm during surgery. Operating rooms are often kept cold for the comfort of the staff, but research shows that for many types of surgery, patients who are kept warm resist infection bet-

ter.⁹³ There are many ways to keep patients warm, including special blankets, hats and booties, and warmed IV liquids.

- **10. Do not shave the surgical site.** Razors can create small nicks in the skin, through which bacteria can enter. If hair must be removed before surgery, ask that clippers be used instead of a razor. ⁹⁴
- 11. Ask that your surgeon limit the number of personnel in the operating room. Every increase in the number of people adds to your risk of infection.⁹⁵
- 12. Ask your doctor about monitoring your glucose (sugar) levels continuously during and after surgery, especially if you are having cardiac surgery. The stress of surgery often makes glucose levels spike erratically. New research shows that when blood glucose levels are tightly controlled to stay between 80–110 mg/unit, heart patients resist infection better. Continue monitoring even when you are discharged from the hospital, because you are not fully healed yet.⁹⁶
- **13.** Avoid a urinary tract catheter if possible. It is a common cause of infection. The tube allows urine to flow from your bladder out of your body. Sometimes catheters are used when busy hospital staff don't have time to walk patients to the bathroom. Ask for a diaper or bed pan instead. They're safer.⁹⁷

- 14. If you must have an IV in your arm, make sure that it is inserted and removed under clean conditions and changed every 3 to 4 days. Intravenous catheters, or IVs, are a common source of infection and are not always necessary. If you need one, insist that it be inserted and removed under clean conditions, which means that your skin is cleaned at the site of insertion, and the person treating you is wearing clean gloves. Alert hospital staff immediately if any redness appears.
- 15. Wash your hands frequently, avoid touching your hands to your mouth, and do not set food or utensils on furniture or bed sheets. Germs such as "C.Diff" can live for many days on surfaces and can cause infections if they get in your mouth.

Note to expectant mothers: If you are planning to have your baby by Cesarean section, follow the steps listed above as if you were having any other type of surgery. Most mothers-to-be probably aren't worried about hospital infections, but if you're having a cesarean, you are ten times more at risk of infection than if you are giving birth vaginally.⁹⁸

The Importance of Hospital Infection Report Cards

Maureen Daly wishes she had known more when she took her 63-year old mother to the hospital. Johanna had slipped and broken her shoulder at a restaurant, and no one expected that she would be in the hospital for more than a day or two. But a *Staph* infection ravaged her body for four months and killed her. "What happened to my mother shouldn't happen to anyone," says Daly. "If only I had had enough information to choose a hospital with a better infection record."

If you need to be hospitalized, wouldn't you want to know which hospital in your area has the lowest infection rate? Good luck getting that information!

The federal Centers for Disease Control and Prevention collect infection data from several hundred hospitals around the nation, but the CDC also promises hospitals to keep infection rates secret. 99 Government, for the most part, is not helping you choose a safe hospital.

The irony is that it's easy to get information for the less important decisions you make in life, such as where to have lunch. Most states will help you find out which restaurants and delicatessens have been cited for health violations. But you can't find out which hospital has the worst infection rate. You can go home to make your own sandwich, but you can't perform surgery on yourself.

The good news is that Colorado, Connecticut, Delaware, Florida. Minnesota, Illinois, Maryland, Missouri, New Hampshire, New Jersey, New York, Ohio, Oklahoma, Oregon, Pennsylvania, Rhode Island, South Carolina, Tennessee, Texas, Virginia, Vermont, and Washington have passed laws to provide the public with hospital infection report cards. Publicly comparing hospital performance will motivate hospitals to improve.

New York's experience with another type of hospital report card proves this. In 1989, New York became the first state to publish each hospital's risk-adjusted mortality rate for cardiac bypass surgery. The results? Deaths from bypass surgery dropped 40 percent, giving New York the lowest mortality rate in the nation for that procedure. Oritics of hospital report cards speculate that deaths went down in New York because hospitals avoided treating the sickest patients, fearing that high-risk

operations would bring down the hospital's grade. However, the evidence proves that's untrue. Deaths declined for a different reason: hospitals forced their worst-performing surgeons — generally, those with low volume - to stop doing the procedure. Patients of the 27 barred surgeons were more than three times as likely to die during surgery. In technical jargon, the 27 surgeons had an average risk-adjusted mortality rate of 11.9 percent, compared with a statewide average of 3.1 percent.¹⁰¹ Wisconsin also found that report cards motivate poorly performing hospitals to improve, according to a 2001 study of 24 hospitals there. 102

Is there a reason *not* to have infection report cards? The hospital industry argues that publicly comparing hospital infection rates would be unfair to hospitals that treat AIDS, cancer, and organ transplant patients who are especially vulnerable to infection. Fair enough, but reports can be risk-adjusted to reflect these differences. What is unfair is keeping the public uninformed.

Fortunately, several other states are considering legislation to provide the public with the information they need. These states should use the model bill suggested here (Appendix A), because it improves upon the laws already passed in three ways: First, it specifies the method of risk-adjustment for surgical site infections used by the CDC, rather than leaving the risk-adjustment method to be determined by committee. This should assure hospitals that comparisons will be fair and take into account which hospitals treat especially sick

and infection-prone patients. 103

Secondly, the bill imposes civil penalties on hospitals that fail to report or flagrantly underreport their infections. These penalties are needed. For many years, some hospitals have openly ignored data collection laws with impunity. For example, in one recent year, hospitals in New York reported only 16.5 percent of the post-surgical deaths that the law required them to report. 104 In 2005, the first year of Pennsylvania's hospital infection reporting program, hospitals reported only one tenth as many infections to the new program as they billed. Some Pennsylvania hospitals implausibly claimed they had no infections at all.105

Thirdly, the model bill ensures that hospital infection reporting will benefit the public, not enrich trial lawyers. The bill provides that "none of the data collected and reported under this law can be used in litigation against an individual hospital."

Next time you hear an ad on the radio urging you to use a particular hospital because it has the best doctors or the latest equipment, keep in mind what you're not being told: how many patients get infections while in that hospital. Hospitals are doing their best to keep that information secret. In contrast, in England hospital infection rates are posted conspicuously on the front door of the hospital. Americans deserve the same information. The legislation proposed here won't help hospitals save face, but it will help you choose a safe hospital. Let hospitals vie for your business by improving their infection rates.

Appendix A

RID's Model Hospital Infection Report Card Bill

T he following outline is intended to help state lawmakers as they draft legislation to provide the public with hospital infection rates:

AN ACT to provide the public with information on infection rates at hospitals in the state of _____.

Section 1. Definitions.

- (a) The public health law is amended to add a new section (lawmakers here should include the specific title of the public health or health department law to be amended).
- (b) "Hospital" shall mean (lawmakers here should consider whether to include only acute care hospitals or also free-standing outpatient surgical centers).
- (c) "Hospital-acquired infection" shall mean, as defined by the federal Centers for Disease Control and Prevention (CDC), "any localized or systemic condition resulting from an adverse reaction to the presence of an infectious agent(s) or its toxin(s) that (a) occurs in a patient in a hospital, (b) and was found not to be present or incubating at the time of admission to the hospital, unless (c) the infection was related to a previous admission to the same hospital."
- (d) "Risk adjustment" shall mean a statistical procedure for comparing patient outcomes, taking into account the differences in patient populations, including risk factors such as the number of patients on central line catheters, or the number of patients undergoing specific types of surgery, as a percentage of the overall number of patients treated. For purposes of this bill, risk adjustment shall duplicate the CDC's NNIS System surgical wound infection risk index or use the number of central-catheter days as a risk-adjustment factor for central line infections.

Appendix A

Section 2.

- (a) Using established public health surveillance methods, each hospital shall maintain a program of identifying and tracking the following types of hospital-acquired infections for the purpose of reporting such data semi-annually to the state health department (lawmakers insert the appropriate state department here): central line-associated, laboratory confirmed primary bloodstream infections contracted by intensive care unit patients, and surgical site infections.¹⁰⁶
- (b) The state health department (lawmakers insert the appropriate department name here) shall establish an advisory committee that includes recognized experts in the field of hospital-acquired infection, public reporting of hospital data, and health care quality management to establish data collection and analysis methodologies and risk adjustment procedures.
- (c) The state health department (lawmakers insert the appropriate department name here) shall establish a state-wide data base of all risk-adjusted, hospital-specific infection rates and make it available to the public on a web-site and in printed materials that can be used by consumers, purchasers of healthcare and advocacy groups to compare the performance of individual hospitals, and the aggregate performance of hospitals in the state with those in other states and nationwide.
- (d) The first year of data submission under this section shall be considered the "pilot phase" of the reporting system. The pilot phase is to ensure the completeness and accuracy of hospital reporting and the fairness and completeness of the state health department's report to the public. During this pilot phase, hospital identifiers shall be encrypted, the state health department (lawmakers insert proper department name here) shall provide each hospital with an encryption key for that hospital only, and no public hospital comparisons will be available. Sixty days after the end of the second year of data submission, the state health department (appropriate department name here) will provide its first report to the public with hospital-specific infection rates included.
- (e) To ensure compliance with this law and the accuracy of self-reporting by the hospitals, the department shall establish an audit process. A civil penalty of \$_____ shall be imposed on any hospital that fails to report on time, or is shown to substantially underreport infections, for each semi-annual reporting period.
- (f) None of the data collected and reported under this law can be used in litigation against an individual hospital.

Appendix B

A Model Hospital Screening Bill

The following outline is intended to help state lawmakers as they draft legislation to mandate MRSA screening in acute care hospitals:

AN ACT to provide universal screening for MRSA in acute care hospitals in the state of

- (a) Whereas at least one out of every twenty patients contracts an infection in the hospital, and these infections kill more people in the U.S. each year as all the cases of AIDS, all breast cancer cases, and all auto accidents combined.¹⁰⁷
- (b) Whereas the danger is worsening, because increasingly these infections cannot be cured with commonly used antibiotics. They are "drug-resistant."
- (c) Whereas in 1974, only 2% of Staphylococcus aureus infections were drug resistant, by 1995 that figure had soared to 22%, by 2003, 64%, and now an even higher percentage are drug resistant or MRSA (methicillin-resistant Staphylococcus aureus).
- (d) Whereas hospitals in Holland, Denmark and Finland once faced similarly soaring rates of drug-resistant infections and brought them down below 1% through meticulous hygiene and screening all patients for the bacterium MRSA.¹⁰⁸
- (e) Whereas an increasing number of leading medical centers in the U.S. have proved screening is essential and effective here too, including University of Pittsburgh-Presbyterian, which reduced MRSA infections 90% in an intensive care unit; The Veterans Hospital in Pittsburgh, which reduced MRSA infections 85% in a pilot program, Evanston-Northwestern, Johns Hopkins in Baltimore, St. Raphael's in New Haven, Connecticut, Brigham & Women's in Boston,¹⁰⁹ and numerous other examples.
- (f) Whereas research shows that you cannot prevent MRSA infections until you identify which patients bring these bacteria into the hospital.¹¹⁰ Patients who unknowingly carry MRSA on their skin and in their nasal passages shed it in tiny particles on bedrails, blood pressure cuffs, stethoscopes, furniture and the floor, where the bacteria can live for many days. Patients become infected with MRSA when the bacteria enter their body, usually via a urinary tract catheter, IV catheter, surgical incision or other open wound, or a ventilator.

Appendix B

- (g) Whereas MRSA testing is simple, noninvasive, and no more costly than a routine HIV test now recommended for all patients admitted to hospitals.
- (h) Whereas research indicates that hospital infections add at least \$30.5 billion a year to the nation's health tab in hospital costs alone, including at least \$2 billion in New York State.¹¹¹
- (i) Whereas research demonstrates that in Pennsylvania, in 2005, the average charge for patients who developed a hospital infection (\$173,206) was nearly four times as high as for patients with the same diagnosis and severity of illness who did not contract an infection (\$44,367),and that hospital infections added over two billion dollars in hospital charges in that state alone.¹¹²
- (j) Whereas "virtually all published analysis that have compared the cost of screening of patients on admission and using contact precautions with colonized patients with the cost savings made by preventing healthcare associated MRSA infections have concluded that the combination of surveillance cultures and barrier precautions results in cost savings for hospitals," according to the prestigious medical journal Lancet, and whereas such research proves that "the costs of caring for patients who become infected with MRSA are much greater than the costs of screening programs."¹¹³
- (k) By January 1, 2009, all acute care hospitals in ______ shall screen all patients undergoing in-patient orthopedic or cardiac surgery and all patients entering intensive care units, burn units, and other "high risk units" for the bacterium MRSA. Patients in these categories who are scheduled for hospitalization can be tested up to two weeks before admission. In addition to other best practices, hospitals shall require contact (barrier) precautions when treating patients who test positive for MRSA colonization, including wearing gloves and gowns for all direct patient contact, using dedicated or disposable equipment such as wheelchairs, blood pressure cuffs, and EKG wires, and thoroughly cleaning other equipment after contact with MRSA positive patients. Wherever possible, hospitals will isolate or cohort patients colonized or infected with MRSA, control and monitor their movement within the hospital, and take whatever steps are needed to stop the transmission of MRSA bacteria to patients who did not come into the hospital with it.
- (I) Patients being discharged from intensive care and other high risk units shall be tested again for MRSA, and those testing positive will be informed of their status, and it will be noted in their medical records.
- (m) An acute care hospital that is in violation of the provisions of this Act shall be subject to such penalties as the Department of Health may determine, and in addition, shall be ineligible for Medicaid reimbursement until the violation is corrected.
- (n) This law shall expire on January 1, 2012.

Society for Healthcare Epidemiologists of America Guideline for Preventing Nosocomial Transmission of Multidrug-Resistant Strains of *Staphylococcus Aureus* and *Entrococcus*

Strength of Recommendations				
Category Type	Category Subtype	Recommendation		
Ι	A	Strongly recommended for implementation and strongly supported by well-designed experimental, clinical, or epidemiologic studies.		
	В	Strongly recommended for implementation and supported by some experimental, clinical, or epidemiologic studies and a strong theoretical rationale.		
	С	Required for implementation, as mandated by federal regulation, state regulation, or both or standard.		
II		Suggested for implementation and supported by suggestive clinical or epidemiologic studies or a theoretical rationale.		
No recommendation		Unresolved issue. Practices for which insufficient evidence or no consensus regarding efficacy exists.		

Recommendations

I. Active Surveillance Cultures to Identify the Reservoir for Spread

- **1.** Implement a program of active surveillance cultures and contact precautions to control the spread of epidemiologically significant antibiotic-resistant pathogens known to be spreading in the healthcare system via direct and indirect contact.(IA)29,30,43,45-47,49,57,96,99,102,106,119,138-147,149,171-173,176
- **2.** Surveillance cultures are indicated at the time of hospital admission for patients at high risk for carriage of MRSA, VRE, or both. (IB)71,76,177,320,321
- **3.** Periodic (eg, weekly) surveillance cultures are indicated for patients remaining in the hospital at high risk for carriage of MRSA, VRE, or both because of ward location, antibiotic therapy, underlying disease, duration of stay, or all four. (IA)30,57,102, 137,141,147-149,174,181

- **4.** In facilities found to have a high prevalence on initial sampling, a facility-wide culture survey is indicated to identify all colonized patients and allow implementation of contact precautions. (IB)102,145,322
- **5.** Because transmission occurs throughout the healthcare system, these measures should be implemented in all types of healthcare facilities throughout the system. (IB)119,161,176,182,323
- **6.** The frequency of active surveillance cultures should be based on the prevalence of the pathogen and risk factors for colonization. For example, more frequent cultures are needed in a facility where 50% of all S. aureus isolates are MRSA than in one where less than 1% of all S. aureus isolates are MRSA. (IB)29,30,43,45-47,49,57,96,99, 102,106,119,138-147,149,171-173,176
- **7.** The goal of this program should be to identify every colonized patient, so that all colonized patients are cared for in contact (or cohort) isolation to minimize spread to other patients. (IB)29,30,43,45-47,49,57,96,99,102,106,119,138-147,149,171-173,176
- **8.** Surveillance cultures for VRE should use stool samples or swab samples from the rectum or perirectal area. Polymerase chain reaction, culture with broth enhancement, and quantitative stool culture have each been more sensitive than directly plated rectal or perirectal swab cultures, but the latter have been associated with control of infections and can be recommended as effective and cost-effective until less costly methods of using the other procedures become available. (IB)99,102,106,137,149,181
- 9. VRE patients can be routinely cohorted with other VRE patients. (II) 102,106,145
- **10.** Surveillance cultures for MRSA should always include samples from the anterior vestibule of the nose. (IB)78,315,324
- 11. If present, areas of skin breakdown should also be sampled for MRSA. (IB)315,324
- 12. Throat cultures have been shown to detect S. aureus and MRSA with sensitivity equal to or greater than that of nasal cultures in multiple patient populations. If used, the throat swab can be plated onto the same agar as the nasal swab. This would enhance sensitivity without adding the cost of an extra culture. (IB)67,74
- **13.** Perirectal–perineal cultures have been shown to detect MRSA with high sensitivity in certain patient populations, but the perirectal–perineal area should not be selected as the only site for culture. (IB)315,324,325
- $\bf 14.$ Patients colonized or infected with MRSA isolates can be cohorted with other MRSA patients. (II) $30,\!43,\!45$

- 15. Patients with MRSA isolates that are eradicable because of known susceptibility to multiple drugs useful for eradication (eg, mupirocin, rifampin, minocycline, trimethoprim-sulfamethoxazole, or all four) should not be cohorted with those with isolates resistant to these drugs, if eradication will be used as an adjunctive measure. (II)272
- **16.** In certain settings, such as nursing homes and psychiatric wards, identification of colonized patients is important, but contact precautions may require modification allowing for social contact while limiting physical contact. (II) 119,182,323

II. Hand Hygiene

- 1. HCWs should be encouraged to decontaminate (clean) their hands with an antiseptic-containing preparation before and after all patient contacts. (IA)121,326-330
- 2. Soap and water hand washing is required when hands are visibly dirty or visibly contaminated with blood, body fluids, or body substances. (IA)331
- **3.** When hands are not visibly contaminated with blood, body fluids, or body substances, use of an alcohol hand rub containing an emollient should be encouraged. (IB)215,332-338
- **4.** Lotion compatible with (ie, that does not inactivate) the antiseptic being used should be provided for use by HCWs. (II)339-343
- ${f 5.}$ Monitoring of hand hygiene compliance and feedback to HCWs should be done to motivate greater compliance. (IB)215,344

III. Barrier Precautions for Patients Known or Suspected to Be Colonized or Infected With Epidemiologically Important Antibiotic-Resistant Pathogens Such as MRSA or VRE

- 1. Gloves should always be worn to enter the room of a patient on contact precautions for colonization or infection with antibiotic-resistant pathogens such as MRSA, VRE, VISA, or VRSA. (IA)122,132,212,225-230
- **2.** Gowns always should be worn as part of contact precautions for all patient and environmental contact with patients known to be colonized by antibiotic-resistant pathogens such as MRSA, VRE, VISA, or VRSA, except when there is no direct contact with patient or environmental surfaces. (IA)29,30,43,4547,49,57,59,96, 99,102,106,119,122,132,135,136,138-147,149,171-173,176,345
- **3.** Universal gown and glove use or universal gloving alone also can be considered for adjunctive control on high-risk wards among patients with surveillance cultures pending. (IB)37,44,105,316-318,346

4. Masks should be worn as part of isolation precautions when entering the room of a patient colonized or infected with MRSA, VISA, or VRSA to decrease nasal acquisition by HCWs. (II)30,123,124,129,231,232

IV. Antibiotic Stewardship

- 1. Avoid inappropriate or excessive antibiotic prophylaxis and therapy. (IB) 194,251,347
- 2. Ensure correct dosage and duration of antibiotic therapy. (IB)348-350
- **3.** Restrict the use of vancomycin (if possible and appropriate for care of the individual patient being treated) to decrease the selective pressure favoring vancomycin resistance. (IB)115,269
- 4. То prevent the establishment of VRE intesticolonization, decrease the use of agents with little activity against enterococci, such as third-generation and fourth-generation cephalosporins, in patients not known to be VRE colonized (if possible and individual patient being appropriate for care of the (IB)115,267,268,351,352
- **5.** To prevent persistent high-density VRE colonization, decrease the use of antianaerobic agents in patients with known VRE intestinal colonization (if possible and appropriate for care of the individual patient being treated). (II) 102,113,159,270
- **6.** To help prevent persistent carriage of MRSA, reduce the use of antibiotics and particularly fluoroquinolones to the minimum necessary in institutions where MRSA is endemic. (IB)251-258
- **7.** Avoid therapy for colonization except when suppression or eradication of colonization is being attempted using an evidence-based approach for infection prevention, for psychological benefit of the patient, or for cost benefit (ie, by reducing the need for long-term isolation). (IB)5,272,285,286

V. Decolonization or Suppression of Colonized Patients

- **1.** Consider MRSA decolonization therapy for both patients and HCWs as an adjunctive measure for controlling spread of MRSA in selected populations when appropriate. (IB)30,176,271,272,275-277
- 2. Any program of decolonization therapy should incorporate routine susceptibility testing, as selection of inactive agents is less likely to achieve eradication. (II)272,353
- **3.** Widespread use, prolonged use, or both of decolonization therapy should be avoided, because this has been associated with the evolution and spread of antibiotic- resistant strains, undermining the effectiveness of the control effort.

(IB)285,286

VI. Other

- 1. Educational programs should be conducted to ensure that HCWs understand why antibiotic-resistant pathogens are epidemiologically important, why prevention of spread is critically necessary for control, and which measures for preventing spread have proven effective. (IB)215,220
- **2.** Ensure that the hospital method of disinfecting hospital surfaces for antibiotic-resistant organisms (especially VRE) has been shown to be adequate based on the results of studies of such methods in the healthcare setting or perform cultures in the room of discharged patients to confirm the adequacy of terminal cleaning. This requires review of the disinfectant agent, method and meticulousness of cleaning, dilutions, and contact time. (IB)102,161,169,294
- **3.** Use the hospital computer system to record longterm isolation indicators for patients colonized with MRSA, VRE, VISA, or VRSA so that on return the computer will provide an alert regarding the need for isolation. (IB)297
- **4.** Dedicate the use of noncritical patient-care equipment to a single patient (or cohort of patients infected or colonized with the pathogen requiring precautions) to avoid sharing between patients. If use of common equipment or items is unavoidable, then adequately clean and disinfect them before use for another patient. (IB)99,150-155,296

References

- Centers for Disease Control and Prevention. Campaign to Prevent Antimicrobial Resistance in Healthcare Settings: Why a Campaign? Atlanta, GA: Centers for Disease Control and Prevention; 2001. Available atwww.cdc.gov/drugresistance/healthcare/ problem.htm.
- ² Neu HC. The crisis in antibiotic resistance. Science 1992;257:1064- 1073.
- Marshall G, Crofton JW, Cruickshank R, et al. The treatment of pulmonary tuberculosis with isoniazid. BMJ 1952;2:735-746.
- ⁴ Aubry-Damon H, Soussy CJ, Courvalin P. Characterization of mutations in the rpoB gene that confer rifampin resistance in *Staphylococcus aureus*. *Antimicrob Agents Chemother* 1998;42:2590-2594.
- Schmitz FJ, Fluit AC, Hafner D, et al. Development of resistance to ciprofloxacin, rifampin, and mupirocin in methicillin-susceptible and -resistant *Staphylococcus aureus* isolates. *Antimicrob Agents* Vol. 24 No. 5 PREVENTING SPREAD OF ANTIBIOTIC RESISTANCE 379 *Chemother* 2000;44:3229-3231.
- ⁶ O'Neill AJ, Cove JH, Chopra I. Mutation frequencies for resistance to fusidic acid and rifampin in Staphylococcus aureus. J Antimicrob Chemother 2001;47:647-650.
- Eisenstadt E, Carlton BC, Brown BJ. Gene mutation. In: Gerhardt P, Murray RGE, Wood WA, Krieg NR, eds. *Methods for General and Molecular Bacteriology*. Washington, DC: American Society for Microbiology; 1994:297-316.
- ⁸ Kucers A, Bennett N. The Use of Antibiotics, 4th ed. London: Heinemann Medical Books; 1987.
- 9 Holmes O. The contagiousness of puerperal fever. N Engl Q J Med Surg

- 1842-1843;1:501-530.
- ¹⁰ Semmelweis I. The Etiology, the Concept and the Prophylaxis of Childbed Fever. Pest, Hungary: CA Hartleben's Verlag-Expedition; 1861.
- ¹¹ Darwin C. On the Origin of Species by Means of Natural Selection. London: J. Murry; 1859.
- ¹² Centers for Disease Control and Prevention. National Nosocomial Infections Surveillance (NNIS) System report: data summary from January 1990–May 1999. Am J Infect Control 1999;27:520-532.
- ¹³ Centers for Disease Control and Prevention. NNIS Antimicrobial Resistance Report: Vancomycin-Resistant Enterococcus (VRE) Facts. Atlanta, GA: Centers for Disease Control and Prevention; 1999.
- ¹⁴ Centers for Disease Control and Prevention. National Nosocomial Infection Surveillance (NNIS) System report: data summary from January 1992–June 2001. Am J Infect Control 2001;29:404-421.
- 15 Cosgrove SE, Sakoulas G, Perencevich EN, Schwaber MJ, Karchmer AW, Carmeli Y. Comparison of mortality associated with methicillinresistant and methicillin-susceptible *Staphylococcus aureus* bacteremia: a meta-analysis. *Clin Infect Dis* 2003;36:53-59.
- ¹⁶ Carmeli Y, Cosgrove SE, Harbarth S, Karchmer AW, Kaye KS, Qi Y. The impact of methicillin-resistance in *Staphylococcus aureus* bacteremia on patient outcomes: mortality, length of stay and hospital charge. Presented at the 41st Annual Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; December 16-19, 2001; Chicago, IL. Abstract K-1221:415.
- ¹⁷ Engemann JJ, Carmeli Y, Cosgrove SE, et al. Adverse clinical and economic outcomes attributable to methicillin resistance among patients with *Staphylococcus aureus* surgical site infection. *Clin Infect Dis* 2003;36:592-598.
- ¹⁸ Salgado C, Farr B. The cost of vancomycin-resistance (VR): a metaanalysis. Presented at the 12th Annual Meeting of the Society for Healthcare Epidemiology of America; April 6-9, 2002; Salt Lake City, UT. Abstract 113:67.
- ¹⁹ Benson K, Hartz AJ. A comparison of observational studies and randomized controlled trials. N Engl J Med 2000;342:1878-1886.
- ²⁰ Concato J, Shah N, Horwitz RJ. Randomized controlled trials, observational studies, and the hierarchy of research designs. N Engl J Med 2000;342:1887-1892.
- ²¹ Hiramatsu K. Molecular evolution of MRSA. *Microbiol Immunol* 1995;39:531-543.
- ²² Hiramatsu K, Cui L, Kuroda M, Ito T. The emergence and evolution of methicillin-resistant Staphylococcus aureus. Trends Microbiol 2001;9:486-493.
- ²³ Kreiswirth B, Kornblum J, Arbeit WE, et al. Evidence for a clonal origin of methicillin-resistance in *Staphylococcus aureus*. *Science* 1993;259:227-230.
- ²⁴ Oliveira DC, Tomasz A, de Lencastre H. The evolution of pandemic clones of methicillin-resistant Staphylococcus aureus: identification of two ancestral genetic backgrounds and the associated mec elements. *Microb Drug Resist* 2001;7:349-361.
- ²⁵ Musser J, Kapur V. Clonal analysis of methicillin-resistant *Staphylococcus aureus* from intercontinental sources: association of the mec gene with divergent phylogenetic lineages implies dissemination by horizontal transfer and recombination. *J Clin Microbiol* 1992;30:2058-2063.
- ²⁶ Givney R, Vickery A, Holliday A, Pegler M, Benn R. Evolution of an endemic methicillin-resistant *Staphylococcus aureus* population in an Australian hospital from 1967-1996. *J Clin Microbiol* 1998;36:552-556.
- ²⁷ Crisostomo MI, Westh H, Tomasz A, Chung M, Oliveria DC, de Lencastre H. The evolution of methicillin resistance in *Staphylococcus aureus*: similarity of genetic backgrounds in historically early methicillin-susceptible and resistant and contemporary epidemic clones. *Proc Natl Acad Sci U S A* 2001;98:9865-9870.
- ²⁸ Enright MC, Robinson DA, Randle G, Feil DJ, Grundmann H, Spratt BG. The evolutionary history of methicillin-resistant *Staphylococcus aureus* (MRSA). *Proc Natl Acad Sci U S A* 2002;99:7687-7692.
- ²⁹ Haley RW, Cushion NB, Tenover FC, et al. Eradication of endemic methicillin-resistant Staphylococcus aureus infections from a neonatal intensive care unit. J Infect Dis

- 1995;171:614-624.
- Jernigan JA, Titus MG, Groschel DHM, Getchell-White SI, Farr BM. Effectiveness of contact isolation during a hospital outbreak of methicillin- resistant *Staphylococcus aureus*. Am J Epidemiol 1996;143:496-504.
- ³¹ Salmenlinna S, Lyytikainen O, Kotilainen P, Scotford R, Siren E, Vuopio-Varkila J. Molecular epidemiology of methicillin-resistant *Staphylococcus aureus* in Finland. *Eur J Clin Microbiol Infect Dis* 2000;19:101-107.
- ³² Roberts RB, de Lancastre A, Eisner W, et al. Molecular epidemiology of methicillin-resistant Staphylococcus aureus in 12 New York hospitals: MRSA collaborative group. J Infect Dis 1998;178:164-171.
- ³³ de Lancastre H, Severina EP, Roberts RB, Kreiswirth B, Tomasz A. Testing the efficacy of a molecular surveillance network: methicillinresistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant *Enterococcus faecium* (VREF) genotypes in six hospitals in the metropolitan New York City area. The BARG Initiative Pilot Study Group. Bacterial Antibiotic Resistance Group. *Microb Drug Resist* 1996;2:343-351.
- ³⁴ Roman RS, Smith J, Walker M, et al. Rapid geographic spread of a methicillin-resistant *Staphylococcus aureus* strain. *Clin Infect Dis* 1997;25:698-705.
- 35 Villari P, Farullo C, Torre I, Nani E. Molecular characterization of methicillin-resistant Staphylococcus aureus (MRSA) in a university hospital in Italy. Eur J Epidemiol 1998;14:802-816.
- ³⁶ Diekema DJ, Pfaller MA, Turnidge J, et al. Genetic relatedness of multidrug- resistant, methicillin-resistant *Staphylococcus aureus* bloodstream isolates: SENTRY antimicrobial resistance surveillance centers worldwide. *Microb Drug Resist* 2000;6:213-221.
- ³⁷ Vriens MR, Fluit AC, Troelstra A, Verhoef J, Van Der Werken C. Are MRSA more contagious than MSSA in a surgical intensive care unit. *Infect Control Hosp Epidemiol* 2002;23:491-494.
- ³⁸ Witte W, Cuny C, Braulke C, Heuck D, Klare I. Widespread dissemination of epidemic MRSA in German hospitals. *Eurosurveillance* 1997;2:25-28.
- ³⁹ Austin DJ, Anderson RM. Transmission dynamics of epidemic methicillin- resistant Staphylococcus aureus and vancomycin-resistant enterococci in England and Wales. J Infect Dis 1999;179:883-891.
- ⁴⁰ Deplano A, Witte W, Van Leeuwen WJ, Brun Y, Struelens MJ. Clonal dissemination of epidemic methicillin-resistant *Staphylococcus aureus* in Belgium and neighboring countries. *Clin Microbiol Infect* 2000;6:239-245.
- ⁴¹ Galdbart JO, Morvan A, El Solh N. Phenotypic and molecular typing of nosocomial methicillin-resistant *Staphylococcus aureus* strains susceptible to gentamicin isolated from France from 1995-1997. *J Clin Microbiol* 2000; 8:185-190.
- ⁴² Sanchez IS, Ramirez M, Troni H, et al. Evidence for geographic spread of a methicillin-resistant *Staphylococcus aureus* clone between Portugal and Spain. *J Clin Microbiol* 1995;33:1234-1236.
- ⁴³ Thompson RL, Cabezudo I, Wenzel RP. Epidemiology of nosocomial infections caused by methicillin-resistant Staphylococcus aureus. Ann Intern Med 1982;97:309-317.
- ⁴⁴ Verhoef J, Beaujean D, Blok H, et al. A Dutch approach to methicillinresistant Staphylococcus aureus. Eur J Clin Microbiol Infect Dis 1999; 18:461-466.
- ⁴⁵ Jernigan JA, Clemence MA, Stott GA, et al. Control of methicillin resistant *Staphylococcus aureus* at a university hospital: one decade later. *Infect Control Hosp Epidemiol* 1995;16:686-696.
- ⁴⁶ Chaix C, Durand-Zaleski I, Alberti C, Brun-Buisson C. Control of endemic methicillin-resistant *Staphylococcus aureus*: a cost benefit analysis in an intensive care unit. *JAMA* 1999;282:1745-1751.
- ⁴⁷ Jans B, Suetens C, Struelens M. Decreasing MRSA rates in Belgian hospitals: results from the national surveillance network after introduction of national guidelines. Infect Control Hosp Epidemiol 2000;21:419.
- ⁴⁸ Bager F. DANMAP 98: Consumption of Antimicrobial Agents and Occurrence of Antimicrobials in Bacteria From Food Animals, Food and Humans in Denmark. Copenhagen: Statens Serum Institut, Danish Veterinary and Food Administration,

- Danish Medicines Agency, Danish Veterinary Laboratory; 1998. Available at www.svs.dk/dk/z/Danmap%201998.pd. 1999.
- ⁴⁹ Harbarth S, Martin Y, Rohner P, Henry N, Auckenthaler R, Pittet D. Effect of delayed infection control measures on a hospital outbreak of methicillin-resistant *Staphylococcus aureus*. *J Hosp Infect* 2000;46:43-49.
- Merrer J, Santoli F, Appere de Vecchi C, Tran B, De Jonghe B, Outin H. "Colonization pressure" and risk of acquisition of methicillin-resistant *Staphylococcus aureus* in a medical intensive care unit. *Infect Control Hosp Epidemiol* 2000;21:718-723.
- ⁵¹ Farr BM, Salgado CD, Karchmer TB, Sherertz RJ. Can antibiotic-resistant nosocomial infections be controlled? *Lancet Infect Dis* 2001;1:38-45.
- ⁵² Vincent JL, Bihari DJ, Suter PM, et al. The prevalence of nosocomial 380 INFECTION CONTROL AND HOSPITAL EPIDEMIOLOGY May 2003 infection in intensive care units in Europe: results of the European Prevalence of Infection in Intensive Care (EPIC) Study. EPIC International Advisory Committee. *JAMA* 1995;274:639-644.
- ⁵³ Cars O, Molstad S, Melander A. Variation in antibiotic use in the European Union. *Lancet* 2001;357:1851-1853.
- ⁵⁴ Frank MO, Batteiger BE, Sorensen SJ, et al. Decrease in expenditures and selected nosocomial infections following implementation of an antimicrobial-prescribing improvement program. *Clinical Performance and Quality Healthcare* 1997;5:180-188.
- Fukatsu K, Saito HK, Matsuda T, Ikeda S, Furukawa S, Muto T. Influences of type and duration of antimicrobial prophylaxis on an outbreak of methicillin-resistant *Staphylococcus aureus* and on the incidence of wound infection. *Arch Surg* 1997;132:1320-1325.
- ⁵⁶ Batteiger BE. Personal communication. Indianapolis: Indiana University; 2001.
- ⁵⁷ Back NA, Linnemann CC Jr, Staneck JL, Kotagal UR. Control of methicillin-resistant Staphylococcus aureus in a neonatal intensive-care unit: use of intensive microbiologic surveillance and mupirocin. Infect Control Hosp Epidemiol 1996;17:227-231.
- ⁵⁸ Barrett FF, McGehee RF, Finland M. Methicillin-resistant Staphylococcus aureus at Boston City Hospital. N Engl J Med 1968;279:441-448.
- ⁵⁹ Boyce JM. Are the epidemiology and microbiology of methicillin-resistant *Staphylococcus aureus* changing? JAMA 1998;279:623-624.
- ⁶⁰ Brumfitt W, Hamilton-Miller J. Methicillin-resistant Staphylococcus aureus. N Engl J Med 1989;320:1188-1196.
- ⁶¹ Naimi TS, LeDell KH, Boxrud D, et al. Epidemiology and clonality of community-acquired methicillin-resistant *Staphylococcus aureus* in Minnesota, 1996-1998. *Clin Infect Dis* 2001;33:990-996.
- ⁶² Frank AI, Marcinak JF, Mangat PD, Schreckeberger PC. Communityacquired and clindamycin-susceptible methicillin-resistant *Staphylococcus aureus* in children. *Pediatr Infect Dis J* 1999;18:993- 1000.
- ⁶³ Suggs AH, Maranan MC, Boyle-Vavra S, Daum RS. Methicillin-resistant and borderline methicillin-resistant asymptomatic *Staphylococcus* colonization in children without identifiable risk factors. *Pediatr Infect Dis J* 1999;18:410-414.
- ⁶⁴ Herold BD, Immergluck LC, Maranan MC, et al. Community-acquired methicillin in children with no identified predisposing risk. *IAMA* 1998;279:593-598.
- ⁶⁵ Adcock PM, Pastor P, Medley F, Patterson JE, Murphy TV. Methicillinresistant *Staphylococcus aureus* (MRSA) in two child care centers. *J Infect Dis* 1998;178:577-580.
- ⁶⁶ Embil J, Ramotar K, Romance L, et al. Methicillin-resistant *Staphylococcus aureus* in tertiary care institutions on the Canadian prairies 1990-1992. Infect Control Hosp Epidemiol 1994;15:646-651.
- ⁶⁷ Shahin R, Johnson IL, Jamieson F, McGeer A, Tolkin J, Ford-Jones EL. Methicillin-resistant Staphylococcus aureus carriage in a child care center following a case of disease. Arch Pediatr Adolesc Med 1999;153:864-868.
- ⁶⁸ Centers for Disease Control and Prevention. Four pediatric deaths from community-acquired methicillin-resistant *Staphylococcus aureus*: Minnesota and North Dakota, 1997-1999. *MMWR* 1999;48:707-710.
- ⁶⁹ Morin CA, Hadler JL. Population-based incidence and characteristics of community-onset

- Staphylococcus aureus infections with bacteremia in 4 metropolitan Connecticut areas, 1998. J Infect Dis 2001;184:1029-1034.
- ⁷⁰ Layton MC, Hierholzer WJ, Patterson JE. The evolving epidemiology of methicillin-resistant Staphylococcus aureus at a university hospital. Infect Control Hosp Epidemiol 1995;16:12-17.
- 71 Troillet N, Carmeli Y, Samore MH, et al. Carriage of methicillin-resistant Staphylococcus aureus at hospital admission. Infect Control Hosp Epidemiol 1998;19:181-185.
- ⁷² Groom AV, Wolsey DH, Naimi TS, et al. Community-acquired methicillin- resistant *Staphylococcus aureus* in a rural American Indian community. *JAMA* 2001;286:1201-1205.
- ⁷³ Salmenlinna S, Lyytikainen O, Vuopio-Varkila J. Community-acquired methicillin-resistant Staphylococcus aureus, Finland. Emerg Infect Dis 2002;8:602-607.
- ⁷⁴ Sa-Leao R, Sanches IS, Couto I, Alves CR, de Lencastre H. Low prevalence of methicillin-resistantstrains among *Staphylococcus aureus* colonizing young and healthy members of the community in Portugal. *Microb Drug Resist* 2001;7:237-245.
- Nopsin B, Mathema B, Martinez J, et al. Prevalence of methicillin-resistant and methicillin-susceptible *Staphylococcus aureus* in the community. *J Infect Dis* 2000;182:359-362.
- Muto CA, Cage EG, Durbin LJ, Simonton BM, Farr BM. The utility of culturing patients on admission transferred from other health care facilities for methicillin-resistant *Staphylococcus aureus* (MRSA). Presented at the Ninth Annual Meeting of the Society for Healthcare Epidemiology of America; April 18-20, 1999; San Francisco, CA. Abstract M33:67.
- ⁷⁷ Calfee DP, Durbin LJ, Germanson TP, Toney DM, Smith EB, Farr BM. Prevalence of methicillin-resistant Staphylococcus aureus (MRSA) among household contacts of individuals with nosocomially-acquired MRSA. Infect Control Hosp Epidemiol. 2003. In press.
- Namour MD, Widmer AF, Bale MJ, Jones RN, Wenzel RP. Efficient detection and long-term persistence of the carriage of methicillin-resistant *Staphylococcus aureus*. *Clin Infect Dis* 1994;19:1123-1128.
- ⁷⁹ Gold HS. Vancomycin-resistant enterococci: mechanisms and clinical observations. *Clin Infect Dis* 2001;33:210-219.
- 80 Bonten MJ, Willems R, Weinstein RA. Vancomycin-resistant enterococci: why are they here, and where do they come from? *Lancet Infect Dis* 2001;1:314-325.
- ⁸¹ Murray BE. What can we do about vancomycin-resistant enterococci? Clin Infect Dis 1995;20:1134.
- ⁸² Martone WJ. Spread of vancomycin-resistant enterococci: why did it happen in the United States? *Infect Control Hosp Epidemiol* 1998;19:539-545.
- ⁸⁵ Carmeli Y, Samore MH, Huskins C. The association between antecedent vancomycin treatment and hospital-acquired vancomycinresistant enterococci: a meta-analysis. *Arch Intern Med* 1999;159:2461-2468.
- 84 Salgado C, Calfee DP, Giannetta ET, Farr BM. Rate of turning culture positive for vancomycin-resistant enterococcus after treatment with oral vancomycin. Presented at the 39th Annual Meeting of the Infectious Diseases Society of America; October 25-28, 2001; San Francisco, CA. Abstract (507):126.
- 85 Coque TM, Tomayko JF, Ricke SC, Okhyusen PC, Murray BE. Vancomycin-resistant enterococci from nosocomial, community, and animal sources in the United States. Antimicrob Agents Chemother 1996;40:2605-2609.
- Morris JG, Shay DK, Hebden JN, et al. Enterococci resistant to multiple antimicrobial agents including vancomycin: establishment of endemicity in a university medical center. Ann Intern Med 1995;123:250-259.
- ⁸⁷ Cetinkaya Y, Falk P, Mayhall CG. Vancomycin-resistant enterococci. Clin Microbiol Rev 2000;13:686-707.
- ⁸⁸ Bates J. Epidemiology of vancomycin-resistant enterococcus in the community and the relevance of farm animals to human infection. *J Hosp Infect* 1997;37:89-101.
- 89 Wegener HC, Aarestrup FM, Jensen LB, Hammerum AM, Bager F. Use of antimicrobial

- growth promoters in food animals and Enterococcus faecium resistance to therapeutic antimicrobial drugs in Europe. *Emerg Infect Dis* 1999;5:329-335.
- ⁹⁰ Klare I, Heier H, Claus H, Reissbrodt R, Witt W. vanA-mediated highlevel glycopeptide resistance in Enterococcus faecium from animal husbandry. FEMS Microbiol Lett 1995;125:165.
- ⁹¹ Devriese LA, Ieven M, Goossens H, et al. Presence of vancomycinresistant enterococci in farm and pet animals. *Antimicrob Agents Chemother* 1996;40:2285-2287.
- ⁹² Borgen K, Simonsen G, Sundsfjord A, Wasteson Y, Olsvik O, Kruse H. Continuing high prevalence of VanA-type vancomycin-resistant enterococci on Norwegian poultry farms three years after avoparcin was banned. *J Appl Microbiol* 2000;89:478-485.
- ⁹⁵ Willems R, Homan W, Top J, et al. Variant esp gene as a marker of a distinct genetic lineage of vancomycin-resistant *Enterococcus faecium* spreading in hospitals. *Lancet* 2001;357:853-855.
- ⁹⁴ Rice LB, Carias L, Rudin S, et al. A potential virulence gene, hylEfm, predominates in Enterococcus faecium of clinical origin. J Infect Dis 2003;187:508-512.
- 95 Boyce JM, Mermel LA, Zervos MJ, et al. Controlling vancomycin-resistant enterococci. Infect Control Hosp Epidemiol 1995;16:634-637.
- ⁹⁶ Boyce JM, Opal SM, Chow JW, et al. Outbreak of multi-drug resistant *Enterococcus faecium* with transferable vanB class vancomycin resistance. *J Clin Microbiol* 1994;32:1148-1153.
- ⁹⁷ Clark NC, Cooksey RC, Hill BC, Swenson JM, Tenover FC. Characterization of glycopeptide-resistant enterococci from U.S. hospitals. *Antimicrob Agents Chemother* 1993;37:2311-2317.
- ⁹⁸ Handwerger S, Raucher B, Altarac D, et al. Nosocomial outbreak due to *Enterococcus faecium* highly resistant to vancomycin, penicillin, and gentamicin. *Clin Infect Dis* 1993;16:750.
- ⁹⁹ Livornese LL, Dias S, Romanowski B, et al. Hospital-acquired infection with vancomycin-resistant *Enterococcus faecium* transmitted by electronic thermometers. *Ann Intern Med* 1992;117:112-116.
- ¹⁰⁰ Kim WJ, Weinstein RA, Hayden MK. The changing molecular epidemiology and establishment of endemicity of vancomycin resistance in enterococci at one hospital over a 6-year period. *J Infect Dis* 1999;179:163-171.
- ¹⁰¹ Moreno RM, Grota P, Crisp C, et al. Clinical and molecular epidemiVol. 24 No. 5 PREVENTING SPREAD OF ANTIBIOTIC RESISTANCE 381 ology of vancomycin-resistant Enterococcus faecium during its emergence in a city in Southern Texas. Clin Infect Dis 1995;21:1234-1237.
- ¹⁰² Byers KE, Anglim AM, Anneski CJ, et al. A hospital epidemic of vancomycin-resistant enterococcus: risk factors and control. *Infect Control Hosp Epidemiol* 2001;22:140-147.
- ¹⁰³ Handwerger S, Skoble J, Discottoo LF, Pucci MJ. Heterogeneity of the vanA gene in clinical isolates of enterococci from the Northeastern United States. *Antimicrob Agents Chemother* 1995;39:362-368.
- ¹⁰⁴ Mato R, de Lancesstre H, Carraher M, Robers RB, Tomasz A. Multiplicity of genetic backgrounds among vancomycin-resistant Enterococcus faecium isolates recovered from an outbreak in a New York City Hospital. *Microb Drug Resist* 1996;2:309-317.
- ¹⁰⁵ Slaughter S, Hayden MK, Nathan C, et al. A comparison of the effect of universal use of gloves and gowns with that of glove use alone on acquisition of vancomycin-resistant enterococci in a medical intensive care unit. *Ann Intern Med* 1996;125:448-456.
- ¹⁰⁶ Muto CA, Posey K, Pokrywka M, et al. The value of identifying the vancomycin resistant enterococci (VRE) reservoir using weekly VRE surveillance culturing (VRESC): "the iceberg melts." Presented at the 12th Annual Meeting of the Society for Healthcare Epidemiology of America; April 6-9, 2002; Salt Lake City, UT. Abstract.
- ¹⁰⁷ Nelson RR, McGregor KF, Brown AR, Amyes GS, Young H. Isolation and characterization of glycopeptide-resistant enterococci from hospitalized patients over a 30 month period. *J Clin Microbiol* 2000;38:2112-2116.
- Stosor V, Kruszynski J, Suriano T, Noskin GA, Peterson LR. Molecular epidemiology of vancomycin-resistant enterococci: a 2-year perspective. Infect Control Hosp Epidemiol

- 1999;20:653-659.
- ¹⁰⁹ de Lencastre H, Brown AE, Chung M, Armstrong D, Tomasz A. Role of transposon Tn5482 in the epidemiology of vancomycin-resistant *Enterococcus faecium* in the pediatric oncology unit of a New York City hospital. *Microb Drug Resist* 1999;5:113-129.
- ¹¹⁰ Beezhold DW, Slaughter S, Hayden MK, et al. Skin colonization with vancomycin-resistant enterococci among hospitalized patients with bacteremia. *Clin Infect Dis* 1997;24:704-706.
- ¹¹¹ Tornieporth NG, Roberts RB, John J, Hafner A, Riley LW. Risk factors associated with vancomycin-resistant *Enterococcus faecium* infection or colonization in 145 matched case patients and control patients. *Clin Infect Dis* 1996;23:767-772.
- ¹¹² Bonten MJ, Hayden MK, Nathan C, et al. Epidemiology of colonization of patients and environment with vancomycin-resistant enterococci. *Lancet* 1996;348:1615-1619.
- ¹¹³ Donskey CJ, Chowdhry T, Hecker M, et al. Effect of antibiotic therapy on the density of vancomycin-resistant enterococci in the stool of colonized patients. *N Engl J Med* 2000;343:1925-1932.
- ¹¹⁴ Bonten MJ, Slaughter S, Ambergen AW, et al. The role of "colonization pressure" in the spread of vancomycin-resistant enterococci: an important infection control variable. *Arch Intern Med* 1998;158:1127-1132.
- ¹¹⁵ Quale J, Landman D, Saurina G, Atwood E, DiTore V, Patel K. Manipulation of a hospital antimicrobial formulary to control an outbreak of vancomycin-resistant enterococci. *Clin Infect Dis* 1996;23:1020-1025.
- ¹¹⁶ Goetz AM, Rihs JD, Wagener MM, Muder RR. Infection and colonization with vancomycin-resistant *Enterococcus faecium* in an acute care Veterans Affairs Medical Center: a 2-year survey. *Am J Infect Control* 1998;26:558-562.
- ¹¹⁷ Fridkin SK, Steward CD, Edwards JR, et al. Surveillance of antimicrobial use and antimicrobial resistance in United States hospitals: Project ICARE phase 2. *Clin Infect Dis* 1999;29:245-252.
- ¹¹⁸ Lautenbach E, LaRosa LA, Marr AM, Nachamkin I, Bilker WB, Fishman NO. Changes in the prevalence of vancomycin-resistant enterococci in response to antimicrobial formulary interventions: impact of progressive restrictions on use of vancomycin and third-generation cephalosporins. *Clin Infect Dis* 2003;36:440-446.
- ¹¹⁹ Ostrowsky BE, Trick WE, Sohn AH, et al. Control of vancomycinresistant enterococcus in health care facilities in a region. N Engl J Med 2001;344:1427-1433.
- ¹²⁰ Montecalvo MA, Jarvis WR, Uman J, et al. Costs and savings associated with infection control measures that reduced transmission of vancomycin-resistant enterococci in an endemic setting. *Infect Control Hosp Epidemiol* 2001;22:437-442.
- ¹²¹ Mortimer EA, Lipsitz PJ, Wolinksy E, et al. Transmission of staphylococci between newborns. Am J Dis Child 1962;104:289-295.
- ¹²² Zachary KC, Bayne PS, Morrison V, Ford DS, Silver LC, Hooper DC. Contamination of gowns, gloves, and stethoscopes with vancomycinresistant enterococci. *Infect Control Hosp Epidemiol* 2001;22:560-564.
- 123 Cookson B, Peters B, Webster M, Phillips I, Rahman M, Noble W. Staff carriage of epidemic methicillin-resistant Staphylococcus aureus. J Clin Microbiol 1989;27:1471-1476.
- ¹²⁴ Lacey S, Flaxman D, Scales J, Wilson A. The usefulness of masks in preventing transient carriage of epidemic methicillin-resistant *Staphylococcus aureus* by healthcare workers. *J Hosp Infect* 2001;48:308-311.
- ¹²⁵ Wells CL, Juni BA, Cameron SB, et al. Stool carriage, isolation, and mortality during outbreak of vancomycin-resistant enterococci in hospitalized medical and/or surgical patients. Clin Infect Dis 1995;21:45-50.
- ¹²⁶ Bonilla HF, Zervos MA, Lyons MJ, et al. Colonization with vancomycin- resistant Enterococcus faecium: comparison of a long-term care unit with an acute-care hospital. Infect Control Hosp Epidemiol 1997;18:333-339.
- ¹²⁷ Noskin GA, Stosor V, Cooper I, Peterson L. Recovery of vancomycinresistant enterococci on fingertips and environmental surfaces. *Infect Control Hosp Epidemiol* 1995;16:577-581.
- ¹²⁸ Suh HK, Jeon YH, Song SJ. A molecular epidemiologic study of methicillin- resistant

- Staphylococcus aureus infection in patients undergoing middle ear surgery. Eur Arch Otorbinolaryngol 1998;255:347-351.
- ¹²⁹ Opal SM, Mayer KH, Stenberg MJ, et al. Frequent acquisition of multiple strains of methicillin-resistant *Staphylococcus aureus* by healthcare workers in an endemic hospital environment. *Infect Control Hosp Epidemiol* 1990;11:479-485.
- 130 Devine J, Cooke RP, Wright EP. Is methicillin-resistant *Staphylococcus aureus* (MRSA) contamination of ward-based computer terminals a surrogate marker for nosocomial MRSA transmission and handwashing compliance. *J Hosp Infect* 2001;48:72-75.
- ¹³¹ Layton MC, Perez M, Heald, Patterson JE. An outbreak of mupirocinresistant Staphylococcus aureus on a dermatology ward associated with an environmental reservoir. Infect Control Hosp Epidemiol 1993;14:369-375.
- ¹³² Boyce JM, Potter-Bynoe G, Chenevert C, King T. Environmental contamination due to methicillin-resistant *Staphylococcus aureus*: possible infection control implications. *Infect Control Hosp Epidemiol* 1997;18:622-627.
- ¹³³ Neely AN, Maley MP. Survival of enterococci and staphylococci on hospital fabrics and plastics. J Clin Microbiol 2000;38:724-726.
- ¹³⁴ Boyce JM, Chenevert C. Isolation gowns prevent health care workers (HCWs) from contaminating their clothing, and possibly their hands, with methicillin-resistant *Staphylococcus aureus* (MRSA) and resistant enterococci. Presented at the Eighth Annual Meeting of the Society for Healthcare Epidemiology of America; April 5-7, 1998; Orlando, FL. Abstract S74:52.
- ¹³⁵ Puzniak LA, Leet T, Mayfield J, Kollef M, Mundy LM. To gown or not to gown: the effect on acquisition of vancomycin resistant enterococci. *Clin Infect Dis* 2002;35:18-25.
- ¹³⁶ Srinivasan A, Song X, Bower R, et al. A prospective study to determine whether cover gowns in addition to gloves decrease nosocomial transmission of vancomycin-resistant enterococci in an ICU. *Infect Control Hosp Epidemiol* 2002;23:424-428.
- ¹³⁷ Calfee DP, Farr BM. Infection control and cost control in the era of managed care. *Infect Control Hosp Epidemiol* 2002;23:407-410.
- ¹³⁸ Karanfil IV, Murphy M, Josephson A, et al. A cluster of vancomycinresistant *Enterococcus faecium* in an intensive care unit. *Infect Control Hosp Epidemiol* 1992;13:195-200.
- ¹³⁹ Montecalvo MA, Horowitz H, Gedris C, Carbonaro C, Tenover FC, Issah A. Outbreak of vancomycin-, ampicillin-, and aminoglycosideresistant *Enterococcus faecium* bacteremia in an adult oncology unit. *Antimicrob Agents Chemother* 1994;38:1363-1367.
- ¹⁴⁰ Dembry L, Uzokwe K, Zervos M. Control of endemic glycopeptideresistant enterococci. *Infect Control Hosp Epidemiol* 1996;17:286-292.
- ¹⁴¹ Rupp ME, Marion N, Fey PD, et al. Outbreak of vancomycin-resistant *Enterococcus fae-cium* in a neonatal intensive care unit. *Infect Control Hosp Epidemiol* 2001;22:301-303.
- ¹⁴² Malik RK, Montecalvo MA, Reale MR, et al. Epidemiology and control of vancomycin-resistant enterococci in a regional neonatal intensive care unit. *Pediatric Infect Dis J* 1999;18:352-356.
- ¹⁴³ Muto CA, Karchmer TB, Cage EG, Durbin LJ, Simonton B, Farr BM. The utility of culturing roommates of patients with vancomycin resistant enterocccus. Presented at the Eighth Annual Meeting of the Society for Healthcare Epidemiology of America; April 5-7, 1998; Orlando, FL. Abstract 76:38.
- ¹⁴⁴ Rubin LG, Tucci V, Cercenado E, Elipoulos G, Isenberg HD. Vancomycin-resistant Enterococcus faecium in hospitalized children. Infect Control Hosp Epidemiol 1992;13:700-705.
- ¹⁴⁵ Jochimsen E, Fish L, Manning K, et al. Control of vancomycin-resistant enterococci at a community hospital: efficacy of patient and staff cohorting. *Infect Control Hosp Epidemiol* 1999;20:106-109.
- ¹⁴⁶ Golan Y, Sullivan B, Snydman DR. Elimination of vancomycin-resistant enterococcus (VRE) transmission in a neonatal intensive care unit (NICU). Presented at the 39th Annual Meeting of the Infectious 382 INFECTION CONTROL AND HOSPITAL EPIDEMIOLOGY May 2003 Diseases Society of America; October 25-28, 2001; San Francisco, CA. Abstract 209:75.
- ¹⁴⁷ Price CS, Paule S, Noskin GA, Peterson LR. Active surveillance reduces vancomycin-resistant

- enterococci (VRE) bloodstream isolates. Presented at the 39th Annual Meeting of the Infectious Diseases Society of America; October 25-28, 2001; San Francisco, CA. Abstract 212:75.
- ¹⁴⁸ Siddiqui AH, Harris AD, Hebden J, Wilson PD, Morris JG, Roghmann M. The effect of active surveillance for vancomycin resistant enterococci in high risk units on vancomycin resistant enterococci incidence hospital-wide. *Am J Infect Control* 2002;30:40-43.
- ¹⁴⁹ Calfee DP, Giannetta E, Farr BM. Effective control of VRE colonization using CDC recommendations for detection and isolation. Presented at the 38th Annual Meeting of the Infectious Diseases Society of America; September 7-10, 2000; New Orleans, LA. Abstract 21:44.
- ¹⁵⁰ Ackelsberg J, Kostman J. A laboratory and clinical study of stethoscopes as potential fomites of infection. Presented at the 33rd Annual Meeting of the Infectious Diseases Society of America; September 16-18, 1995; San Francisco, CA.
- ¹⁵¹ Bernard L, Kereveur A, Durand D, et al. Bacterial contamination of hospital physicians' stethoscopes. *Infect Control Hosp Epidemiol* 1999;20:626-628.
- ¹⁵² Breathnach AS, Jenkins DR, Pedler SJ. Stethoscopes as possible vectors of infection by staphylococci. BMJ 1992;305:1573-1574.
- ¹⁵³ Maki DG, Halvorson K, Fisher S. The stethoscope: a medical device with potential for amplifying cross-infection of resistant nosocomial organisms in the hospital. Presented at the 36th Annual Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 15-18, 1996; New Orleans, LA. Abstract 154:247.
- ¹⁵⁴ Smith MA, Mathewson JJ, Ulert IA, Scerpella EG, Ericsson CD. Contaminated stethoscopes revisited. Arch Intern Med 1996;156:82-84.
- ¹⁵⁵ Cohen HA, Amir J, Matalon A, Mayan R, Beni S, Barzilai A. Stethoscopes and otoscopes: a potential vector of infection? *Fam Pract* 1997;14:446-449.
- ¹⁵⁶ Singh D, Kaur H, Gardner WG, Treen LB. Bacterial contamination of hospital pagers. Infect Control Hosp Epidemiol 2002;23:274-276.
- ¹⁵⁷ Brooks S, Khan A, Stoica D, Griffith J. Reduction in vancomycin-resistant *Enterococcus* and *Clostridium difficile* infections following change to tympanic thermometers. *Infect Control Hosp Epidemiol* 1998;19:333-336.
- 158 Centers for Disease Control and Prevention/Healthcare Infection Control Practices Advisory Committee (HICPAC). *Draft Guidelines for Environmental Infection Control* in Healthcare Facilities. Atlanta, GA: Centers for Disease Control and Prevention; 2001. Available at www.cdc.gov/ncidod/hip/enviro/guide.htm.
- 159 Edmond MB, Ober JF, Weinbaum DL, et al. Vancomycin-resistant Enterococcus faecium bacteremia: risk factors for infection. Clin Infect Dis 1995;20:1126-1133.
- ¹⁶⁰ Wendt C, Wiesenthal B, Dietz E, Ruden H. Survival of vancomycinresistant and vancomycin-susceptible enterococci on dry surfaces. J Clin Microbiol 1998;36:3734-3736.
- ¹⁶¹ Smith TL, Iwen PC, Olson SB, Rupp ME. Environmental contamination with vancomycin-resistant enterococci in an outpatient setting. *Infect Control Hosp Epidemiol* 1998;19:515-518.
- 162 Oie S, Kamiya A. Survival of methicillin-resistant Staphylococcus aureus (MRSA) on naturally contaminated dry mop. J Hosp Infect 1996;34:145-149.
- ¹⁶³ Noskin GA, Peterson L, Warren J. Enterococcus faecium and Enterococcus faecalis bacteremia: acquisition and outcome. Clin Infect Dis 1995;20:296-301.
- ¹⁶⁴ Dietz B, Raht A, Wendt C, Martiny H. Survival of MRSA on sterile goods packaging. J Hosp Infect 2001;49:255-261.
- ¹⁶⁵ McDade J, Hall L. Survival of Staphylococcus aureus in the environment: I. Exposure on surfaces. American Journal of Hygiene 1963;78:330-337.
- ¹⁶⁶ Rutala W, Katz E, Sherertz R, Sarubbi F. Environmental study of methicillin-resistant Staphylococcus aureus epidemic in a burn unit. J Clin Microbiol 1983;18:683-688.
- ¹⁶⁷ Embil J, McLeod J, Al-Barrak AM, et al. An outbreak of methicillinresistant *Staphylococcus aureus* on a burn unit: potential role of contaminated hydrotherapy equipment. *Burn* 2001;27:681-688.
- 168 Noskin GA, Bednarz P, Suriano T, Reiner S, Peterson LR. Persistent contamination of fabric-covered furniture by vancomycin-resistant enterococci: implications for uphol-

- stery selection in hospitals. Am J Infect Control 2000;28:311-313.
- ¹⁶⁹ Falk PS, Winnike J, Woodmansee C, Desai M, Mayhall CG. Outbreak due to vancomycin-resistant enterococci (VRE) in a burn unit. Infect Control Hosp Epidemiol 2000;21:575-582.
- ¹⁷⁰ Rimland D. Nosocomial infections with methicillin and tobramycin resistant Staphylococcus aureus: implication of physiotherapy in hospital-wide dissemination. Am J Med Sci 1985;290:91-97.
- ¹⁷¹ Law MR, Gill ON. Hospital-acquired infection with methicillin-resistant and methicillin-sensitive staphylococci. *Epidemiol Infect* 1988;101:623-629.
- ¹⁷² Murray-Leisure KA, Geib S, et al. Control of epidemic methicillinresistant *Staphylococcus aureus. Infect Control Hosp Epidemiol* 1990;11:343-350.
- ¹⁷³ Nicolle LE, Dyck B, Thompson G, et al. Regional dissemination and control of epidemic methicillin-resistant *Staphylococcus aureus*. *Infect Control Hosp Epidemiol* 1999;20:202-205.
- 174 Cantey J, Rhoton B, Southgate W, Snyder C. Control of spread of methicillin resistant Staphylococcus aureus in a neonatal ICU. Presented at the 12th Annual Meeting of the Society for Healthcare Epidemiology of America; April 6-9, 2002; Salt Lake City, UT. Abstract 36:49.
- ¹⁷⁵ Croyle K, Muto C. Surveillance cultures in the race against MRSA: ahead by a nose. Presented at the 12th Annual Meeting of the Society for Healthcare Epidemiology of America; April 6-9, 2002; Salt Lake City, UT. Abstract 35:49.
- ¹⁷⁶ Kotilainen P, Routamaa M, Peltonen R, et al. Eradication of methicillin-resistant Staphylococcus aureus from a health center ward and associated nursing home. Arch Intern Med 2001;161:859-863.
- ¹⁷⁷ Nouer A, Araujo A, Chebabo A, Cardoso F, Pinto M, Hospital Universitário Universidade Federal do Rio de Janeiro. Control of methicillin-resistant *Staphylococcus aureus* (MRSA) in an intensive care unit (ICU) after the institution of routine screening. Presented at the 42nd General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA. Abstract K-97:97.
- ¹⁷⁸ Horcajada J, Marco F, Martinez J, et al. Prevalence of methicillinresistant *Staphylococcus aureus* colonization at admission in a tertiary hospital: usefulness of early detection. Presented at the 42nd General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA. Abstract K-98.
- ¹⁷⁹ Gerard M, Dediste A, Van Esse R, et al. Cost effectiveness of a policy of methicillin resistant *Staphylococcus aureus* (MRSA) screening, decontamination and isolation in a medical ICU. Presented at the 42nd General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA. Abstract K-99.
- ¹⁸⁰ Green K, Fleming CA, Richardson H, Low DE, Willey B, McGeer A. MRSA and VRE in Ontario, Canada: results of 7 years of surveillance and control measures. Presented at the 42nd General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA. Abstract K661.
- ¹⁸¹ Muto CA, Giannetta ET, Durbin LJ, Simonton BM, Farr BM. Cost effectiveness of perirectal surveillance cultures for controlling vancomycin-resistant enterococcus. *Infect Control Hosp Epidemiol* 2002;23:429-435.
- 182 Armstrong-Evans M, Litt M, Willey B, et al. Control of transmission of vancomycin-resistant Enterococcus faecium in a long-term-care facility. Infect Control Hosp Epidemiol 1999;20:312-317.
- ¹⁸³ Hill AB. A Short Textbook of Medical Statistics, vol. 11. London: Hodder and Stoughton; 1984.
- ¹⁸⁴ Salgado C, Sherertz R, Karchmer T, et al. Public health initiative to control MRSA and VRE in Virginia and North Carolina. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 164:75.
- ¹⁸⁵ Arnold MS, Dempsey JM, Fishman M, McAuley PJ, Tibert C, Vallande NC. The best hospital practices for controlling methicillinresistant Staphylococcus aureus: on the culturing

- edge. Infect Control Hosp Epidemiol 2002;23:69-76.
- ¹⁸⁶ D'Agata EMC, Thayer V, Shaffner W. An outbreak of Acinetobacter baumannii: the importance of cross transmission. Infect Control Hosp Epidemiol 2000;21:588-591.
- ¹⁸⁷ Simor AE, Lee M, Vearncombe M, et al. An outbreak due to multiresistant Acinetobacter baumannii in an acute burn care unit: risk factors for acquisition and management. Infect Control Hosp Epidemiol 2002;23:261-267.
- Piagnerelli M, Carlier E, Deplano A, Lejeune P, Govaerts D. Risk factors for infection and molecular typing in patients in the intensive care unit colonized with nosocomial Enterobacter aerogenes. Infect Control Hosp Epidemiol 2002;23:452-456.
- ¹⁸⁹ Selkon JB, Stokes ER, Ingham HR. The role of an isolation unit in the control of hospital infection with methicillin resistant staphylococci. *J Hosp Infect* 1980;1:41-46.
- ¹⁹⁰ Pearman JW, Christiansen KJ, Annear DI, et al. Control of methicillin resistant Staphylococcus aureus (MRSA) in an Australian metropolitan Vol. 24 No. 5 PREVENTING SPREAD OF ANTIBIOTIC RESISTANCE 383 teaching hospital complex. Med J Aust 1985;142:103-108.
- ¹⁹¹ Shanson DC, Johnstone D, Midgley J. Control of a hospital outbreak of methicillin-resistant *Staphylococcus aureus* infections: value of an isolation unit. *J Hosp Infect* 1985;6:285-292.
- ¹⁹² Price EH, Brain A, Dickson JAS. An outbreak with a gentamicin and methicillin-resistant Staphylococcus aureus in a neonatal unit. J Hosp Infect 1980;1:221-228.
- ¹⁹³ Farrington M, Redpath C, Trundle C, Coomber S, Brown NM. Winning the battle but losing the war: methicillin-resistant *Staphylococcus aureus* (MRSA) infection at a teaching hospital. *QIM* 1998;91:539-548.
- ¹⁹⁴ Kunin CM, Tupasi T, Craig WA. Use of antibiotics: a brief exposition of the problem and some tentative solutions. *Ann Intern Med* 1973;79:555-560.
- ¹⁹⁵ Zuckerman RA, Steele L, Venezia RA, Tobin EH. Undetected vancomycin-resistant Enterococcus in surgical intensive care unit patients. Infect Control Hosp Epidemiol 1999;20:685-686.
- ¹⁹⁶ Ostrowsky B, Venkataraman L, D'Agata E, Gold H, DeGirolami P, Samore M. Vancomycin-resistant enterococci in intensive care units: high frequency of stool carriage during a non-outbreak period. *Arch Intern Med* 1999;159:1467-1472.
- ¹⁹⁷ Austin DJ, Bonten MJM, Weinstein RA, Slaughter S, Anderson RM. Vancomycin-resistant enterococci in intensive-care hospital settings: transmission dynamics, persistence, and the impact of infection control programs. *Proc Natl Acad Sci U S A* 1999;96:6908-6913.
- ¹⁹⁸ Sebile V, Chevret S, Valleron A. Modeling the spread of resistant nosocomial pathogens in an intensive care unit. Infect Control Hosp Epidemiol 1997;18:84-92.
- ¹⁹⁹ Warren DK, Kollef MH, Seiler SM, Fridken SK, Fraser VJ. The epidemiology of vancomycin-resistant Enterococcus colonization in a medical intensive care unit. *Infect Control Hosp Epidemiol* 2003;24:257-263.
- ²⁰⁰ Ridwan B, Mascini E, van Der RN, Verhoef J, Bonten M. What action should be taken to prevent spread of vancomycin resistant enterococci in European hospitals? *BMJ* 2002;324:666-668.
- ²⁰¹ Brisse S, Fussing V, Ridwan B, Verhoef J, Willems RJ. Automated ribotyping of vancomycin resistant *Enterococcus faecium* isolates. *J Clin Microbiol* 2002;40:1977-1984.
- ²⁰² Francois P, Pittet D, Bento M, et al. Rapid detection of methicillinresistant *Staphylococcus aureus* directly from sterile or nonsterile clinical samples by a new molecular assay. *J Clin Microbiol* 2003;41:254-260.
- ²⁰⁵ Lankford MG, Zembower TR, Trick WE, Hacek DM, Noskin GA, Peterson LR. Impact of hospital design on the handwashing compliance among healthcare workers. Presented at the 38th Annual Meeting of the Infectious Diseases Society of America; September 7- 10, 2000; New Orleans, LA. Abstract 18:43.
- ²⁰⁴ Preston GA, Larson EL, Stamm W. The effect of private isolation rooms on patient care practices, colonization and infection in an intensive care unit. Am J Med 1981;70:641-645.
- ²⁰⁵ Albert RK, Condie F. Handwashing patterns in medical intensive care units. N Engl J Med 1981;304:1465.
- ²⁰⁶ Larson E. Compliance with isolation technique. Am J Infect Control 1983;11:221-225.

- ²⁰⁷ Donowitz LG, Hunt EH, Pugh VG, Farr BM, Hendley JO. Comparison of historical and serologic immunity to varicella-zoster virus in 373 hospital employees. *Am J Infect Control* 1987;15:212-214.
- ²⁰⁸ Graham M. Frequency and duration of handwashing in an intensive care unit. Am J Infect Control 1990;18:77-81.
- ²⁰⁹ Dubbert PM, Dolce J, Richter W, Miller M, Chapman S. Increasing ICU staff hand-washing: effects of education and group feedback. *Infect Control Hosp Epidemiol* 1990;11:191-193.
- ²¹⁰ Pettinger A, Nettleman MD. Epidemiology of isolation precautions. *Infect Control Hosp Epidemiol* 1991;12:303-307.
- ²¹¹ Larson EL, McGinley K, Foglia A, et al. Handwashing practices and resistance and density of bacterial hand flora on two pediatric units in Lima, Peru. *Infect Control Hosp Epidemiol* 1992;20:65-72.
- ²¹² Doebbeling BN, Pfaller MA, Houston AK, Wenzel RP. Removal of nosocomial pathogens from the contaminated glove: implications for glove reuse and handwashing. *Ann Intern Med* 1988; 109:394-398.
- ²¹³ Zimakoff J, Kjelsberg AB, Larsen SO, Holstein B. A multicenter questionnaire investigation of attitudes toward hand hygiene, assessed by the staff in fifteen hospitals in Denmark and Norway. *Am J Infect Control* 1992;20:58-64.
- ²¹⁴ Meengs MR, Giles BK, Chisholm CD, Cordell WH, Nelson DR. Hand washing frequency in an emergency department. *J Emerg Nurs* 1994;20:183-188.
- ²¹⁵ Pittet D, Hugonnet S, Harbarth S, et al. Effectiveness of a hospitalwide programme to improve compliance with hand hygiene: infection control programme. *Lancet* 2000;356:1307-1312.
- ²¹⁶ Pittet D, Mourouga P, Perneger TV. Compliance with handwashing in a teaching hospital. Ann Intern Med 1999;130:126-130.
- ²¹⁷ Vernon MO, Trick WE, Welbel SF, Peterson BJ, Weinstein RA. Hand hygiene adherence: does the number of sinks matter? *Infect Control Hosp Epidemiol* 2003;24:224-225.
- ²¹⁸ Boyce JM, Pittet D. Guideline for hand hygiene in health-care settings: recommendations of the Healthcare Infection Control Practices Advisory Committee and the HICPAC/SHEA/APIC/IDSA Hand Hygiene Task Force. MMWR 2002;51(RR16):1-45.
- ²¹⁹ Grundmann H, Hori S, Winter B, Tami A, Austin DJ. Risk factors for the transmission of methicillin-resistant *Staphylococcus aureus* in an adult intensive care unit: fitting a model to the data. *J Infect Dis* 2002;185:481-488.
- ²²⁰ Larson EL, Early E, Cloonan P, Sugrue S, Parides M. An organizational climate intervention associated with increased handwashing and decreased nosocomial infections. *Behav Med* 2000;26:14-22.
- ²²¹ Webster J, Faoagali JL, Cartwright D. Elimination of methicillin-resistant Staphylococcus aureus from a neonatal intensive care unit after hand washing with triclosan. J Paediatr Child Health 1994;30:59-64.
- ²²² Zafar AB, Butler RC, Reese DJ, Gaydos LA. Use of 0.3% triclosan (Bacti-Stat*) to eradicate an outbreak of methicillin-resistant *Staphylococcus aureus* in a neonatal nursery. *Am J Infect Control* 1995;23:200-208.
- ²²³ Lucet JC, Rigaud MP, Mentre F, et al. Hand contamination before and after different hand hygiene techniques: a randomized clinical trial. J Hosp Infect 2002;50:276-280.
- ²²⁴ Pittet D, Dharan S, Touveneau S, Sauvan V, Perneger TV. Bacterial contamination of the hands of hospital staff during routine patient care. *Arch Intern Med* 1999;159:821-826.
- ²²⁵ Klein BS, Perloff WH, Maki DG. Reduction of nosocomial infection during pediatric intensive care by protective isolation. N Engl J Med 1989;320:1714-1721.
- ²²⁶ Leclair JM, Freeman J, Sullivan BF, Crowley CM, Goldmann DA. Prevention of nosocomial RSV infections through compliance with glove and gown isolation precautions. *N Engl J Med* 1987;317:329-334.
- ²²⁷ Olsen RJ, Lynch P, Coyle MB, Cummings J, Bokete T, Stamm WE. Examination gloves as barriers to hand contamination in clinical practice. *JAMA* 1993;270:350-353.
- 228 Tenorio AR, Badri SM, Sahgal NB, et al. Effectiveness of gloves in the prevention of hand carriage of vancomycin-resistant enterococcus species by health care workers after

- patient care. Clin Infect Dis 2001;32:826-829.
- ²²⁹ Johnson S, Gerding DN, Olson MM, et al. Prospective, controlled study of vinyl glove use to interrupt *Clostridium difficile* nosocomial transmission. *Am J Med* 1990;88:137-140.
- ²³⁰ Muto CA, Byers KE, Karchmer TB, Dill JB, Durbin LJ, Farr BM. Controlling vancomycin-resistant enterococcus (VRE) at a university hospital. Presented at the Seventh Annual Meeting of the Society for Healthcare Epidemiology of America; April 27-29, 1997; St. Louis, MO. Abstract 52:28.
- ²³¹ Karchmer TB, Ribadeneyra MG, Durbin LJ, Giannetta E, Farr BM. Prevalence of methicillin-resistant *Staphylococcus aureus* colonization among resident-physicians at a teaching hospital using contact/droplet precautions for MRSA isolation. Presented at the Tenth Annual Meeting of the Society for Healthcare and Epidemiology of America; March 5-9, 2000; Atlanta, GA. Abstract 200.
- ²³² Sherertz RJ, Reagan DR, Hampton KD, et al. A cloud adult: the Staphylococcus aureusvirus interaction revisited. Ann Intern Med 1996;124:539-547.
- ²³³ Williams RE. Epidemiology of airborne staphylococcal infection. *Bacteriol Rev* 1966;30:660-674.
- ²³⁴ Mermel L, Dempsey J, Parenteau S. An MRSA outbreak in a surgical intensive care unit: possible role of aerosol transmission from opened ventilator tubing. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 17:42.
- ²³⁵ Meester M, Schultsz C, Boeijen-Donkers L, et al. Evidence for airborne transmission of methicillin-resistant S. aureus. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 36:46
- ²³⁶ Bischoff W, Bassetti S, Bassetti-Wyss B, et al. 'The cloud phenomenon' (CP): predictors of *Staphylococcus aureus* (SA) airborne dispersal associated with rhinovirus infection. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 100:60.
- ²³⁷ Shiomori T, Miyamoto H, Makishima K. Significance of airborne transmission of methicillin-resistant Staphylococcus aureus in an otolaryngology- head and neck surgery unit. Arch Otolaryngol Head Neck Surg 2001;127:644-648.
- ²³⁸ Anonymous. Staphylococcus aureus resistant to vancomycin: United States, 2002.
 MMWR 2002;51:565-567. 384 INFECTION CONTROL AND HOSPITAL EPIDEMIOLOGY May 2003
- ²³⁹ Achong M, Hauser BA, Krusky JL. Rational and irrational use of antibiotics in a Canadian teaching hospital. *Canadian Medical Association Journal* 1977;116:256.
- ²⁴⁰ Kunin CM. The responsibility of the infectious disease community for the optimal use of antimicrobial agents. *J Infect Dis* 1985;151:388.
- ²⁴¹ Maki DG, Schuna AA. A study of antimicrobial misuse in a university hospital. Am J Med Sci 1978;275:271.
- ²⁴² Roberts AW, Visconti JA. The rational and irrational use of systemic antimicrobial drugs. American Journal of Hospital Pharmacy 1972;29:828.
- ²⁴³ Scheckler WE, Bennett JV. Antibiotic use in seven community hospitals. *JAMA* 1970;214:264.
- ²⁴⁴ Bamberger DM, Dahl SL. Impact of voluntary vs enforced compliance of third generation cephalosporin use in a teaching hospital. *Arch Intern Med* 1992;152:554-557.
- ²⁴⁵ Shah SS, Sinkowitz-Cochran, Keyserling H, Jarvis WR. Vancomycin use in pediatric cardiothoracic surgery patients. *Pediatr Infect Dis J* 1999;18:558-560.
- ²⁴⁶ Shah SS, Sinkowitz-Cochran, Keyserling H, Jarvis WR. Vancomycin use in pediatric neurosurgery patients. *Am J Infect Control* 1999;27:482-497.
- ²⁴⁷ Lawton RM, Fridkin SK, Gaynes RP, McGowan JE. Practices to improve antimicrobial use at 47 hospitals: the status of the 1997 SHEA/IDSA position paper recommendations. *Infect Control Hosp Epidemiol* 2000;21:256-259.
- ²⁴⁸ Hopkins HA, Sinkowitz-Cochran RL, Rudin BA, Keyserling HL, Jarvis WR. Vancomycin use in pediatric hematology-oncology patients. *Infect Control Hosp Epidemiol* 2000;21:48-50.

- ²⁴⁹ Shales DM, Gerding DN, John JF, et al. Society for Healthcare Epidemiology of America and Infectious Diseases Society of America Joint Committee on the Prevention of Antimicrobial Resistance: guidelines for the prevention of antimicrobial resistance in hospitals. *Clin Infect Dis* 1997;25:584-599.
- ²⁵⁰ Forrest A, Nix DE, Ballow CH, Goss TF, Birmingham MC, Schentag JJ. Pharmacodynamics of intravenous ciprofloxacin in seriously ill patients. *Antimicrob Agents Chemother* 1993;37:1073-1081.
- ²⁵¹ Monnet DL. Methicillin-resistant Staphylococcus aureus and its relationship to antimicrobial use: possible implications for control. Infect Control Hosp Epidemiol 1998;19:552-559.
- ²⁵² Weber SG, Gold HS, Karchmer AW, Hooper DC, Carmeli Y. Exposure to quinolones is a risk factor for methicillin-resistant (MRSA), but not for methicillin-susceptible *Staphylococcus aureus* (MSSA). Presented at the 42nd Annual General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA. Abstract K-2070.
- ²⁵³ Dziekan G, Hahn A, Thune K, et al. Methicillin-resistant *Staphylococcus aureus* in a teaching hospital: investigation of nosocomial transmission using a matched case-control study. J Hosp Infect 2000;46:263-270.
- ²⁵⁴ Harbarth S, Harris AD, Carmeli Y, Samore MH. Parallel analysis of individual and aggregated data on antibiotic exposure and resistance in gram-negative bacilli. *Clin Infect Dis* 2001;33:1462-1468.
- ²⁵⁵ Hori S, Sunley R, Tami A, Grundmann H. The Nottingham *Staphylococcus aureus* population study: prevalence of MRSA among elderly in a university hospital. *J Hosp Infect* 2002;50:25-19.
- ²⁵⁶ Campillo B, Dupeyron C, Richardet JP. Epidemiology of hospitalacquired infections in cirrhotic patients: effect of carriage of methicillin- resistant *Staphylococcus aureus* and influence of previous antibiotic therapy and norfloxacin prophylaxis. *Epidemiol Infect* 2001;127:443-450.
- ²⁵⁷ Gruson D, Hilbert G, Vargas F, et al. Rotation and restricted use of antibiotics in a medical intensive care unit: impact on the incidence of ventilator-associated pneumonia caused by antibiotic-resistant gramnegative bacteria. Am J Respir Crit Care Med 2000;162:837-843.
- ²⁵⁸ Landman D, Chockalingam M, Quale JM. Reduction in the incidence of methicillin-resistant Staphylococcus aureus and ceftazidime-resistant Klebsiella pneumoniae following changes in a hospital antibiotic formulary. Clin Infect Dis 1999;28:1062-1066.
- ²⁵⁹ Van der Auwera P, Pensart N, Korten V, Murray BE, Leclercq R. Influence of oral glycopeptide on the fecal flora of human volunteers: selection of highly glycopeptide-resistant enterococci. *J Infect Dis* 1996;173:1129-1136.
- ²⁶⁰ Tucci V, Haran MA, Isenberg H. Epidemiology and control of vancomycin-resistant enterococci in an adult and children's hospital. Am J Infect Control 1997;25:371-376.
- ²⁶¹ Uttley A, Collins C, Naidoo J, George R. Vancomycin-resistant enterococci. *Lancet* 1988;1:57-58.
- ²⁶² Anglim AM, Klym B, Byers KE, Farr BM. Effect of a vancomycin restriction policy on ordering practices during an outbreak of vancomycin-resistant *Enterococcus faecium*. *Arch Intern Med* 1997;157:1132-1136.
- ²⁶³ Shay DK, Maloney SA, Montecalvo M, et al. Epidemiology and mortality risk of vancomycin-resistant enterococcal bloodstream infections. *J Infect Dis* 1995;172: 993-1000.
- ²⁶⁴ Weinstein JW, Roe M, Towns M, et al. Resistant enterococci: a prospective study of prevalence, incidence, and factors associated with colonization in a university hospital. *Infect Control Hosp Epidemiol* 1996;17:36-41.
- ²⁶⁵ Carmeli Y, Eliopoulos GM, Samore MH. Antecedent treatment with different antibiotics as a risk for vancomycin resistant enterococcus. *Emerg Infect Dis* 2002;8:802-807.
- ²⁶⁶ Lucas GM, Lechtzin N, Puryear DW, Yau LL, Moore RD. Vancomycinresistant and vancomycin-susceptible enterococcal bacteremia: comparison of clinical features and outcomes. *Clin Infect Dis* 1998;26:1127-1133.

- ²⁶⁷ Donskey CJ, Hanrahan JA, Hutton RA, Rice LB. Effect of parenteral antibiotic administration on the establishment of colonization with vancomycin- resistant *Enterococcus faecium* in the mouse gastrointestinal tract. *J Infect Dis* 2000;181:1830-1833.
- ²⁶⁸ Bradley SJ, Wilson ALT, Allen MC, Sher HA, Goldstone AH, Scott GM. The control of hyperendemic glycopeptide-resistant *Enterococcus* spp. on a haematology unit by changing antibiotic usage. *J Antimicrob Chemother* 1999;43:261-266.
- ²⁶⁹ Fridkin SK, Lawton R, Edwards JR, Tenover FC, McGowan JE, Gaynes RP. Monitoring antimicrobial use and resistance: comparison with a national benchmark on reducing vancomycin use and vancomycin-resistant enterococci. *Emerg Infect Dis* 2002;8:7.
- ²⁷⁰ Donskey CJ, Hanrahan JA, Hutton RA, Rice LB. Effect of parenteral antibiotic administration on persistence of vancomycin-resistant *Enterococcus faecium* in the mouse gastrointestinal tract. *J Infect Dis* 1999;180:384-390.
- ²⁷¹ Struelens MJ, Ronveaux O, Jans B, Mertens R, the Groupement pour le Depistage, 'Etude et la Prevention des Infections Hospitalieres. Methicillin-resistant *Staphylococcus aureus* epidemiology and control in Belgian hospitals, 1991 to 1995. *Infect Control Hosp Epidemiol* 1996;17:503-508.
- ²⁷² Karchmer TB, Jernigan JA, Durbin BM, Simonton BM, Farr BM. Eradication of methicillin-resistant *S. aureus* (MRSA) colonization with different regimens. Presented at the Ninth Annual Meeting of the Society for Healthcare Epidemiology of America; April 18-20, 1999; San Francisco, CA. Abstract 65:42.
- ²⁷³ Bartzokas CA, Paton JH, Gibson MF, Graham R, McLoughlin GA, Croton RS. Control and eradication of methicillin-resistant *Staphylococcus aureus* on a surgical unit. *N Engl J Med* 1984;10:255-259.
- ²⁷⁴ Tuffnell DJ, Croton RS, Hemingway DM, Hartley MN, Wake PN, Garvey RJP. Methicillin-resistant *Staphylococcus aureus*: the role of antisepsis in the control of an outbreak. *J Hosp Infect* 1987;10:255-259.
- ²⁷⁵ Boyce JM, Opal SM, Potter-Bynoe G, Mederios AA. Spread of methicillin- resistant *Staphylococcus aureus* in a hospital and after exposure to a health-care worker with chronic sinusitis. *Clin Infect Dis* 1993;17:496-504.
- ²⁷⁶ Kluytmans J, van Leeuwen W, Goessens W, et al. Food-initiated outbreak of methicillin-resistant *Staphylococcus aureus* analyzed by phenoand genotyping. *J Clin Microbiol* 1995;33:1121-1128.
- ²⁷⁷ Lessing MP, Jordens JZ, Bowlwer IC. When should health care workers be screened for methicillin resistant *Staphylococcus aureus? J Hosp Infect* 1997;35:320-321.
- ²⁷⁸ Doebbeling BN, Breneman DL, Neu HC, et al. Elimination of *Staphylococcus aureus* nasal carriage in health care workers: analysis of six clinical trials with mupirocin calcium ointment. The Mupirocin Collaborative Study Group. *Clin Infect Dis* 1993;17:466-474.
- ²⁷⁹ Doebbeling BN, Regan DR, Pfaller MA, Houston AK, Hollis RJ, Wenzel RP. Long term efficacy of intranasal mupirocin ointment: a prospective cohort study of *Staphylococcus aureus* carriage. *Arch Intern Med* 1994;154:1505-1508.
- ²⁸⁰ Regan DR, Doebbeling BN, Pfaller MA, et al. Elimination of coincident *Staphylococcus aureus* nasal and hand carriage with intranasal application of mupirocin ointment: a prospective cohort study of Staphylococcus aureus carriage. *Ann Intern Med* 1991:114:101-106.
- ²⁸¹ Perl TM, Cullen JJ, Wenzel RP, et al. Intranasal mupirocin to prevent postoperative Staphylococcus aureus infections. N Engl J Med 2002;13:1871-1877.
- ²⁸² Farr BM. Mupirocin to prevent *S. aureus* infections. *N Engl J Med* 2002;13:1905-1906.
- ²⁸³ Harbarth S, Dharan S, Liassine N, Herrault P, Auckenthaler R, Pittet D. Randomized, placebo-controlled, double-blind trial to evaluate the efficacy of mupirocin for eradicating carriage of methicillin-resistant *Staphylococcus aureus*. *Antimicrob Agents Chemother* 1999;43:1412-1416.
- ²⁸⁴ Harbarth S, Dharan S, Liassine N, Herrault P, Auckenthaler R, Pittet Vol. 24 No. 5 PREVENTING SPREAD OF ANTIBIOTIC RESISTANCE 385 D. Risk factors for persistent carriage of methicillin-resistant Staphylococcus aureus. Clin Infect Dis 2000;31:1380-1385.
- ²⁸⁵ Vasquez JE, Walker ES, Franzus BW, Overbay BK, Regan DR, Sarubbi FA. The epidemiology of mupirocin resistance among methicillin-resistant Staphylococcus aureus at a

Appendi<u>x C</u>

- Veterans' Affairs hospital. Infect Control Hosp Epidemiol 2000;21:459-464.
- ²⁸⁶ Miller MA, Dascal A, Portnory J, Medelson J. Development of mupirocin resistance among methicillin-resistant Staphylococcus aureus after widespread use of nasal mupirocin ointment. *Infect Control Hosp Epidemiol* 1996;17:811-813.
- ²⁸⁷ Centers for Disease Control and Prevention. Recommendations for preventing the spread of vancomycin resistance: recommendations of the Hospital Infection Control Practices Advisory Committee (HICPAC). MMWR 1995;44:1-13.
- ²⁸⁸ Centers for Disease Control and Prevention. Guidelines for handwashing and hospital environmental control. *MMWR* 1998;36(suppl):2S.
- ²⁸⁹ Spaulding EH. Chemical disinfection of medical and surgical materials. In: Lawrence CA, Block SS, eds. Disinfection, *Sterilization, and Preservation. Philadelphia: Lea & Febiger*; 1968:517-531.
- ²⁹⁰ Rutala W. APIC guidelines for selection and use of disinfectants. Am J Infect Control 1996;24:313-342.
- ²⁹¹ Saurina G, Landman D, Quale J. Activity of disinfectants against vancomycin-resistant Enterococcus faecium. Infect Control Hosp Epidemiol 1998;18:345-347.
- ²⁹² Rutala W, Stiegel M, Sarubbi F, Weber D. Susceptibility of antibioticsusceptible and antibiotic-resistant hospital bacteria to disinfectants. *Infect Control Hosp Epidemiol* 1997;18:417-421.
- ²⁹³ Anderson RL, Carr JH, Bond WW, Favero MS. Susceptibility of vancomycin-resistant enterococci to environmental disinfectants. *Infect Control Hosp Epidemiol* 1997;18: 195-199.
- ²⁹⁴ Byers KE, Durbin LJ, Simonton BM, Anglim AM, Adal KA, Farr BM. Disinfection of hospital rooms contaminated with vancomycin-resistant *Enterococcus faecium*. *Infect Control Hosp Epidemiol* 1998;19:261-264.
- ²⁹⁵ Rampling A, Wiseman S, Davis L, et al. Evidence that hospital hygiene is important in the control of methicillin resistant *Staphylococcus aureus*. *J Hosp Infect* 2001;49:109-116.
- ²⁹⁶ Mangi RJ, Andriole VT. Contaminated stethoscopes: a potential source of nosocomial infections. *Yale J Biol Med* 1972;45:600-604.
- ²⁹⁷ Pittet D, Safran E, Harbarth S, et al. Automatic alerts for methicillinresistant *Staphylococcus aureus* surveillance and control: role of a hospital information system. *Infect Control Hosp Epidemiol* 1996;17:496-502.
- ²⁹⁸ Wakefield DS, Helms CM, Massanari RM, Mori M, Pfaller MA. Cost of nosocomial infection: relative contributions of laboratory, antibiotic, and per diem costs in serious Staphylococcus aureus infections. Am J Infect Control 1988;16:185-192.
- ²⁹⁹ Wakefield DA, Pfaller MA, Hammons GT, Massanari RM. Use of the appropriateness evaluation protocol for estimating the incremental costs associated with nosocomial infections. *Med Care* 1987;25:481-488.
- ³⁰⁰ Arnow PM, Quimosing EM, Beach M. Consequences of intravascular sepsis. Clin Infect Dis 1993;16:778-784.
- ³⁰¹ Rubin RJ, Harrington CA, Poon A, Dietrich K, Greene JA, Moiduddin A. The economic impact of *Staphylococcus aureus* infection in New York City hospitals. *Emerg Infect Dis* 1999;5:9-17.
- 302 Abramson MA, Sexton DJ. Nosocomial methicillin-resistant and methicillin-susceptible Staphylococcus aureus primary bacteremia: at what costs? Infect Control Hosp Epidemiol 1999;20:408-411.
- ³⁰⁸ Cheng AF, French GL. Methicillin-resistant Staphylococcus aureus bacteremia in Hong Kong. J Hosp Infect 1988;12:91-101.
- ³⁰⁴ Stone PW, Larson E, Kawar LN. A systematic audit or economic evidence linking nosocomial infections and infection control interventions: 1990-2000. Am J Infect Control 2002;30:145-152.
- ³⁰⁵ Kaye KS, Engemann JJ, Mozaffari E, Carmeli Y. Outcomes related to vancomycin resistant *Enterococcus* and methicillin resistant *Staphylococcus aureus*: a comparison of the two types of control groups. Presented at the 42nd Annual General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 27-30, 2002; San Francisco, CA.

- ³⁰⁶ Stosor V, Peterson L, Postelnick M, Noskin G. Enterococcus faecium bacteremia: does vancomycin resistance make a difference? Arch Intern Med 1998;158:522-527.
- ³⁰⁷ Jernigan JA, Hadziyannis SC. Vancomycin-resistant *Enterococcus faecium* (VRE) bacteremia (B) in severely neutropenic patients. Presented at the 36th Annual General Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy; September 15-18, 1996; New Orleans, LA. Abstract J8:219.
- 308 Edmond MB, Jones RN, Pfaller MA, Wallace SE, Wenzel RP. Multicenter surveillance for nosocomial enterococcal bacteremia: a comparison of vancomycin-sensitive vs vancomycin-resistant cases. Presented at the Sixth Annual Meeting of the Society for Healthcare Epidemiology of America; April 21-23, 1996; Washington, DC. Abstract 17:18.
- 309 Lautenbach E, Bilker WB, Brennan PJ. Enterococcal bacteremia: risk factors for vancomycin resistance and predictors of mortality. *Infect Control Hosp Epidemiol* 1999;20:318-323.
- ³¹⁰ Stroud L, Edwards J, Danzing L, Culver D, Gaynes R. Risk factors for mortality associated with enterococcal bloodstream infections. *Infect Control Hosp Epidemiol* 1996;17:576-580.
- ³¹¹ Newel KA, Millis JM, Arnow PM, et al. Incidence and outcome of infection by vancomycin-resistant enterococcus following orthotopic liver transplantation. *Transplant* 1998;65:439-442.
- ³¹² Bhavnani SM, Drake JA, Forrest A, et al. A nationwide, multicenter, case-control study comparing risk factors, treatment, and outcome for vancomycin-resistant and -susceptible enterococcal bacteremia. *Diagn Microbiol Infect Dis* 2000;36:145-158.
- ³¹³ Song X, Srinivisan A, Plaut D, Perl TM. Effect of nosocomial vancomycin-resistant enterococcal bacteremia on mortality, length of stay, and cost. *Infect Control Hosp Epidemiol* 2003;24:251-256.
- ³¹⁴ Karchmer TB, Cage EG, Durbin LJ, Simonton BM, Farr BM. Cost effectiveness of active surveillance cultures for controlling methicillinresistant S. aureus (MRSA). J Hosp Infect 2002;51:126-132.
- ³¹⁵ Papia G, Louie M, Tralla A, Johnson C, Collins V, Simor AE. Screening high-risk patients for methicillin-resistant *Staphylococcus aureus* on admission to the hospital: is it cost effective? *Infect Control Hosp Epidemiol* 1999;20:473-477.
- ³¹⁶ Bronstein M, Kaye K, Sexton D. Gown utilization as a measure of cost of methicillin resistant *Staphylococcus aureus* screening. Presented at the 12th Annual Meeting of the Society for Healthcare Epidemiology of America; April 6-9, 2002; Salt Lake City, UT. Abstract: 47:51.
- ³¹⁷ Lucet J, Chevret S, Durand-Zaleski I, Chastang Cregnier B. Prevalence and risk factors for carriage of methicillin resistant *Staphylococcus aureus* at admission to the intensive care unit. *Arch Intern Med* 2003;163:181-188.
- ³¹⁸ Karchmer TB, Farr BM. Presumptive isolation on admission of patients transferred from facilities with a high-prevalence of MRSA: a cost analysis. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 38:46.
- ³¹⁹ Fridken SK, Hageman J, McDougal LK, et al. Epidemiological and microbiological characterization of infections caused by *Staphylococcus aureus* with reduced susceptibility to vancomycin: United States 1997- 2001. *Clin Infect Dis* 2003;36:429-439.
- ³²⁰ Calfee DP, Giannetta E, Durbin LJ, Farr BM. The increasing prevalence of MRSA and VRE colonization among patients transferred from primary and secondary health care facilities. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 171.
- ³²¹ Muto CA, Cage E, Durbin LJ, Simonton BM, Farr BM. The utility of culturing patients on admission transferred from other hospitals or nursing homes for vancomycin resistant *Enterococcus* (VRE). Presented at the 35th Annual Meeting of the Infectious Diseases Society of America; November 12-15, 1998; Denver, CO. Abstract.
- 322 Muto CA, Patel-Brown S, Krystofiak S, et al. Vancomycin-resistant enterococci (VRE): hospital-wide point prevalence study and epidemiologic description. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3,

- 2001; Toronto, Ontario, Canada. Abstract.
- ³²³ Silverblatt FJ, Tibert C, Mikolich D, et al. Preventing the spread of vancomycin-resistant enterococci in a long-term care facility. *J Am Geriatr Soc* 2000;48:1211-1215.
- ³²⁴ Salgado C, Sheppe S, Dill J, Durbin L, Farr B. The sensitivity of MRSA follow-up cultures. Presented at the 11th Annual Meeting of the Society for Healthcare Epidemiology of America; April 1-3, 2001; Toronto, Ontario, Canada. Abstract 40.
- ³²⁵ Manian FA, Finkle D, Zack J, Meyer L. Routine screening for methicillin- resistant Staphylococcus aureus among patients newly admitted to an acute rehabilitation unit. Infect Control Hosp Epidemiol 2002;23:516-519.
- 326 Larson E, Killien M. Factors influencing handwashing behavior of patient care personnel. Am J Infect Control 1982;10:93-99.
- ³²⁷ Maki DG. The use of antiseptics for handwashing by medical personnel. J Chemother 1989;1(suppl):3-11.
- ³²⁸ Massanari RM, Hierholzer W Jr. A crossover comparison of antiseptic soaps on nosocomial infection rates in intensive care units. Am J Infect Control 1984;12:247-248.
- ³²⁹ Doebbeling BN, Stanley GL, Sheetz CT, et al. Comparative efficacy of alternative hand-washing agents in reducing nosocomial infections in 386 INFECTION CONTROL AND HOSPITAL EPIDEMIOLOGY May 2003 intensive care units. N Engl J Med 1992;327:88-93.
- 330 Ehrenkranz NJ, Alfonso BC. Failure of bland soap handwash to prevent hand transfer of patient bacteria to urethral catheters. *Infect Control Hosp Epidemiol* 1991;12:654-662.
- ³³¹ Larson E. A casual link between handwashing and risk of infection? Examination of the evidence. *Infect Control Hosp Epidemiol* 1988;9:28-36.
- ³³² Larson EL, Eke PI, Laughon BE. Efficacy of alcohol-based hand rinses under frequent-use conditions. *Antimicrob Agents Chemother* 1986;30:542-544.
- 333 Larson EL, Aiello AE, Bastyr J, et al. Assessment of two hand hygiene regimens for intensive care unit personnel. Crit Care Med 2001;29:944-951.
- 334 Boyce JM. Scientific basis for handwashing with alcohol and other waterless antiseptic agents. In: Rutala WA, ed. *Disinfection, Sterilization and Antisepsis: Principles and Practices in Healthcare Facilities.* Washington, DC: Association for Professionals in Infection Control and Epidemiology; 2001:140-151.
- 335 Widmer AF. Replace handwashing with use of a waterless alcohol hand rub? Clin Infect Dis 2000;31:136-143.
- ³³⁶ Maury E, Alzieu M, Baudel JL, Haram N. Availability of an alcohol solution can improve hand disinfection compliance in an intensive care unit. Am J Respir Crit Care Med 2000;162:324-327.
- ³⁵⁷ Boyce JM, Kelliher S, Vallande N. Skin irritation and dryness associated with two hand hygiene regimens: soap and water handwashing versus hand antisepsis with an alcoholic hand gel. *Infect Control Hosp Epidemol* 2000;21:442-448.
- 338 Bischoff WE, Reynolds TM, Sessler CN, Edmond MD, Wenzel RP. Handwashing compliance by healthcare workers: the impact of introducing an accessible alcohol-based hand antiseptic. Arch Intern Med 2000;160:1017-1021.
- ³³⁹ Walsh B, Blakemore PH, Drubu YJ. The effect of handcream on the antibacterial activity of chlorhexidine gluconate. *J Hosp Infect* 1987;9:30-33.
- 340 Berndt U, Wigger-Alberti W, Gabard B, Elsner P. Efficacy of a barrier cream and its vehicle as protective measures against occupational irritant contact dermatitis. *Contact Dermatitis* 2000:42:77-80.
- ³⁴¹ McCormick RD, Buchman TL, Maki D. Double-blind, randomized trial of scheduled use of a novel barrier cream and an oil-containing lotion for protecting the hands of health care workers. *Am J infect Control* 2000;28:302-310.
- ³⁴² Heeg P. Does hand care ruin hand disinfection? *J Hosp Infect 2001*;48(suppl A):S37-S39.
- ³⁴³ Dharan S, Hugonnet S, Pittet D. Evaluation of interference of a hand care cream with alcohol-based hand disinfection. *Occup Environ Med* 2001;49:81-84.
- ³⁴⁴ Mayer JA, Dubbert PM, Miller M, Burkett PA, Chapman S. Increasing handwashing in an intensive care unit. *Infect Control* 1986;7:259-262.
- ³⁴⁵ Quale J, Landman D, Atwood E, et al. Experience with a hospital-wide outbreak of vancomycin-resistant enterococci. Am J Infect Control 1996;24:372-379.

- ³⁴⁶ Muto CA, Durbin LJ, Alexander CH, Karchmer TB, Farr BM. Frequency of community acquired MRSA at a university hospital. Presented at the 35th Annual Meeting of the Infectious Diseases Society of America; September 13-16, 1997; San Francisco, CA. Abstract 347:135.
- ³⁴⁷ Harbarth S, Samore MH, Lichtenberg D, Carmeli Y. Prolonged antibiotic prophylaxis after cardiovascular surgery and its effect on surgical site infections and antimicrobial resistance. *Circulation* 2000;10:2916-2921.
- ³⁴⁸ Gross R, Kinky DE, Weiner M, Morgan AS, Gibson GA, Fishman NO. A randomized controlled trial of a comprehensive antimicrobial management program. *Infect Control Hosp Epidemiol* 2000;21:87-88.
- ³⁴⁹ Gross R, Morgan AS, Kinky DE, Weiner M, Gibson GA, Fishman NO. Impact of a hospital-based antimicrobial management program on clinical and economic outcomes. *Clin Infect Dis* 2001;33:289-295.
- 350 Evans RS, Pestonik SL, Classen DC, et al. A computer-assisted management program for antibiotics and other anti-infective agents. N Engl J Med 1998;338:232-238.
- 351 May AK, Melton SM, McGwin G, Cross JM, Moser SA, Rue LW. Reduction of vancomycin-resistant enterococcal infections by limitation of broad-spectrum cephalosporin use in a trauma and burn intensive care unit. Shock 2000;14:259-264.
- ³⁵² Smith DW. Decreased antimicrobial resistance after changes in antibiotic use. *Pharmacotherapy* 1999;19:129S-132S.
- 355 Walker ES, Vasquez JE, Dula R, Bullock H, Sarubbi FA. Mupirocinresistant, methicillin-resistant Staphylococcus aureus: does mupirocin remain effective? Infect Control Hosp Epidemiol 2003;24:342-346.

Appendix D

The Institute for Healthcare Improvement's 5,000,000 Lives Campaign

In December, 2006, The Institute for Healthcare Improvement enlisted thousands of hospitals across the country in its 5,000,000 Lives Campaign. The goal is to prevent 5,000,000 incidents of harm to patients by December, 2008. Four of the twelve elements of that campaign call on hospitals to implement procedures proven effective in reducing infection. In support of this laudable effort, included here are excerpts from the IHI campaign. For more complete information, consult the Institute's Web site at: http://www.ihi.org/IHI/Programs/Campaign/Campaign.htm

I. The Four Key Components of Preventing Ventilator-Associated Pneumonia:

- 1. Elevation of the head of the bed to between 30 and 45 degrees
- 2. Daily "sedation vacation" and daily assessment of the readiness to extubate
- 3. Peptic ulcer disease prophylaxis
- 4. Deep venous thrombosis prophylaxis (unless contraindicated)

II. The Four Key Components for Preventing Surgical Site Infections:

- 1. Appropriate use of antibiotics, including administering antibiotics within one hour before surgical incision, selecting an antibiotic consistent with national guidelines, and discontinuing prophylactic antibiotics within 24 hours after surgery
- 2. No shaving. Appropriate hair removal, if necessary, with clippers or a depilatory, but not with a razor
- 3. Monitor and maintain patient's glucose levels after surgery, particularly for cardiovascular surgery patients.
- 4. Keep patients' body temperatures at normal levels during and after surgery, especially colorectal surgery, with warmed IV fluids, warming blankets, hats and booties, and other means.

Appendix D

III. The Five Key Components of Preventing Catheter-Related Bloodstream Infections:

- Appropriate hand hygiene, including cleaning hands before and after palpating catheter insertion sites, before and after inserting, replacing, accessing, repairing or dressing an intravascular catheter, whenever hands are soiled or contaminated, before and after removing gloves, etc.
- 2. Maximal barrier protection meaning wearing a cap, mask, sterile gown, and gloves when placing or assisting in the placement of a central line, and ensuring that the patient is covered head to toe in a sterile drape with one small opening for the site of insertion
- 3. Chlorhexidine skin antisepsis before insertion
- Optimal catheter site selection, with the subclavian vein as the preferred site instead of the jugular or femoral sites for non-tunneled catheters in adult patients
- 5. Daily review of central line necessity to prevent unnecessary, prolonged use

IV. Reduce Methicillin-Resistant Staphylococcus aureus (MRSA) Infections

- 1. Hand hygiene
- 2. Decontamination of the environment and equipment
- 3. Active surveillance cultures (ASCs)
- 4. Contact precautions for infected and colonized patients
- 5. Compliance with Central Venous Catheter and Ventilator Bundle

About the Author

Dr. McCaughey is a health policy expert and former Lt. Governor of New York State. In 2004, she founded the Committee to Reduce Infection Deaths (www.hospitalinfection.org), a nationwide educational campaign to stop hospital-acquired infections. She serves as the CEO and Chairman of the organization. RID has made hospital infections a major public issue, provided compelling evidence that preventing infection improves hospital profitability as well as saving lives, and won legislation in over 20 states for public reporting of infection rates. RID has become synonymous with patient safety and clean hospital care.

Dr. McCaughey's research on how to prevent infection deaths has been featured on Good Morning America, the CBS Morning Show, ABC's 20/20, and many other national programs. She has appeared on Fox News Network's Hannity & Colmes, The O'Reilly Factor, CNN's Talk Back Live, and numerous radio programs.

Dr. McCaughey is the author of over one hundred scholarly and popular articles on health policy, infection, medical innovation, the economics of aging, and Medicare. Her writings have appeared in The New York Times, The Wall Street Journal, New Republic, Policy Review, Forbes Magazine, New York Law Journal, Los Angeles Times, U.S. News & World Report, and many other national publications. Her 1994 analysis of the Clinton health plan in the New Republic won a National Magazine Award for the best article in the nation on public policy. Her article on the dangers of "Dumbing Down Medical Care" won the National Media Award from the American Society of Anesthesiologists. She has been profiled in The New Yorker, The New York Times Magazine, New York Magazine, The Washington Post, and other publications.

From 1994 to 1998, she served as Lt. Governor of New York State. She focused on health issues, and her bills became models for legislation in many states and in Congress.

She has taught at Vassar College and Columbia University, and produced prize-winning studies while at two think tanks, the Manhattan Institute and later the Hudson Institute.

Prior to entering the health policy field, Dr. McCaughey earned a Ph.D. in constitutional history. She is the author of two books on that subject, From Loyalist to Founding Father (Columbia University Press), winner of the Bancroft Dissertation Award, and Government by Choice (Basic Books). She also chaired a national commission on reforming the Electoral College in 1992, wrote its report, Electing the President, and testified before Congress on the subject. In 1989, she served as Guest Curator for the Bicentennial Exhibit and related events at the New York Historical Society.

Endnotes

- Robert Weinstein, "Nosocomial Infection Update," *Emerging Infectious Diseases* 4.3 (1998): 416-20. The infection rate, between 5 and 6 infections per 100 admissions, has been steady for more than 28 years, according to Weinstein's research.
- The Federal Centers for Disease Control and Prevention previously estimated that 90,000 people die annually from infections they contract in U.S. hospitals, but the CDC press confirms that it is "working on a new number." Interview with Nicole Coffin, spokesperson for the CDC, Infectious Diseases Division, October 10, 2002 (404-639-2888). The Chicago Tribune puts the death rate even higher, at 103,000. The Chicago Tribune examined hospital records, court records, and federal and state agency data pertaining to 5810 hospitals to reach its estimate. The CDC based their extrapolations on data voluntarily submitted by 315 U.S. hospitals: "How the Chicago Tribune Analyzed Infection Cases," Chicago Tribune (July 21, 2002).
- ³ Institute of Medicine, "Insuring America's Health: Principles and Recommendations," Academic Emergency Medicine 11.4 (2004): 418-22.
- ⁴ See the "Preventing Infections Makes Hospital More Profitable" section of this booklet.
- Institute of Medicine, "Insuring America's Health: Principles and Recommendations," Academic Emergency Medicine 11.4 (2004): 418-22.
- ⁶ Jarvis, William R, et al. "National prevalence of methicillin-resistant *Staphylococcus aureus* in inpatients at US health care facilities, 2006." *American Journal of Infection Control*. Vol. 35, No. 10. 2007.
- Infectious Disease Society of America, *Bad Bugs, No Drugs* (July, 2004): 3. Data are from the Federal Centers for Disease Control and Prevention National Nosocomial Infections Surveillance (NNIS) system report, data summary from January 1992 to June 2003, issued August 2003.
- ⁸ CA Muto, JA Jernigan, et al., "SHEA Guideline for Preventing Nosocomial Transmission of Multidrug-Resistant Strains of Staphylococcus aureus and Enterococcus," *Infection Control* and Hospital Epidemiology 24.5 (2003): 362-86.
- Infectious Disease Society of America, Bad Bugs, No Drugs (July, 2004): Table 3. Data are from the Federal Centers for Disease Control and Prevention National Nosocomial Infections Surveillance (NNIS) system report, data summary from January 1992 to June 2003, issued August 2003.
- 8 CA Muto, JA Jernigan, et al., "SHEA Guideline for Preventing Nosocomial Transmission of Multidrug-Resistant Strains of Staphylococcus aureus and Enterococcus," *Infection Control* and Hospital Epidemiology 24.5 (2003): 362-86.
- 11 Ibid.
- ¹² CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," *Infections in Medicine* 20 (2003): 192-200; RL Thompson et al., "Epidemiology of nosocomial infections caused by *methicillin-resistant Staphylococcus aureus*," *Annals of Internal Medicine* 97.3 (1982): 309-317.
- ¹³ Pittsburgh Regional Healthcare Initiative, PRHI in Progress: 2004 in Review (January 2005): 5.
- ¹⁴ Telephone interview with Dr. Carlene Muto, University of Pittsburgh Medical Center-Presbyterian Hospital, July 15, 2005 (412-629-2566).
- ¹⁵ JM Boyce, "Do Infection Control Measures Work for Methicillin-Resistant Staphylococcus Aureus?" *Infection Control and Hospital Epidemiology* 25.5 (2004): 395-401.
- SS Huang, et al., "Impact of routine intensive care unit surveillance cultures and resultant barrier precautions on hospital-wide methicillin-resistant Staphylococcus aureus bacteremia," *Clinical Infectious Diseases* 43.8 (2006): 971-8.
- BE Ostrowsky et al., "Control of vancomycin-resistant Enterococcus in health care facilities in a region," New England Journal of Medicine 344.19 (2001): 1427-33; see also CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," Infections in Medicine 20 (2003): 192-200
- Hand Hygiene in Healthcare Settings Core, slide 3 of a slide presentation dated June 6, 2003, and currently available on the CDC Web site at http://cdc.gov/handhygiene/down-

- load/hand_hygiene_core.ppt; See also: DM Pittet, "Improving Adherence to Hand Hygiene Practice: a Multidisciplinary Approach," *Emerging Infectious Diseases* 7.2 (2001): 234-40; LC Biant et al., "Eradication of *Methicillin Resistant Staphylococcus Aureus* by 'ring fencing' of elective orthopedic beds," *British Medical Journal* 329.7458 (2004): 149-151.
- 19 SS Huang, et al., "Impact of routine intensive care unit surveillance cultures and resultant barrier precautions on hospital-wide methicillin-resistant Staphylococcus aureus bacteremia," *Clinical Infectious Diseases* 43.8 (2006): 971-8.
- ²⁰ PC Carling et al., "Operating room environmental cleaning an evaluation using a new targeting method," Abstract 280 presented at SHEA's 17th Annual Scientific Meeting (April 2007); PC Carling et al., "Improving patient area cleaning/disinfecting activities in twenty hospitals," Abstract 280 presented at SHEA's 17th Annual Scientific Meeting (April 2007)
- MRSA contaminates patient care rooms 69 percent to 73 percent of the time. See: JM Boyce et al., "Environmental contamination due to methicilliin-resistant Staphylococcus aureus: possible infection control implications," *Infection Control and Hospital Epidemiology* 18.9 (1997): 622-7.
- ²² JM Boyce et al., "Environmental contamination due to methicillin-resistant Staphylococcus aureus: possible infection control implications," Infection Control and Hospital Epidemiology 18.9 (1997): 622-627.
- ²³ Falk PS et al., "Outbreak of Vancomycin-Resistant Enterococci in a Burn Unit," *Infection Control and Hospital Epidemiology* 21.9 (Sept. 2000): 575-582.
- ²⁴ CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," *Infections in Medicine* 20 (2003): 192-200.
- 25 The American Medical Association recommends it. CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," *Infections in Medicine* 20 (2003): 192-200
- ²⁶ AM Treakle et al., "Methicillin-resistant Staphylococcus aureus (MRSA) and white coats of healthcare workers," Abstract 163 presented at SHEA's 17th Annual Scientific Meeting (April 2007).
- ²⁷ JA Jernigan et al., "Effectiveness of Contact Isolation during a Hospital Outbreak of Methicillin-Resistant Staphylococcus aureus," *American Journal of Epidemiology* 143.5 (1996): 496-504.
- ²⁸ Jarvis, William R, et al. "National prevalence of methicillin-resistant *Staphylococcus aureus* in inpatients at US health care facilities, 2006." *American Journal of Infection Control*. Vol. 35, No. 10. 2007.
- ²⁹ Klevens, Monina, et. Al. "Invasive Methicillin-Resistant Staphylococcus aureus Infections in the United States" *Journal of the American Medical Association*. Vol. 298 No. 15, pp. 1733-1826, October 17, 2007.
- ³⁰ S Harbarth, D Pittet, "Control of Nosocomial Methicillin-Resistant Staphylococcus aureus: Where Shall We Send Our Hospital Director Next Time?" *Infection Control and Hospital Epidemiology* 24.5 (2003): 314-6.
- ³¹ BM Farr, "Doing the Right Thing (and Figuring Out What That Is)," *Infection Control and Hospital Epidemiiology* 27.10 (2006): 999-1003, citing data from the CDC National Nosocomial Infection Survey (NNIS).
- ³² DA Goldmann et al., "Strategies to Prevent and Control the Emergence and Spread of Antimicrobial-Resistant Microorganisms in Hospitals. A challenge to hospital leadership," *Journal of the American Medical Association* 275.3 (1996): 234-40.
- ³³ CA Muto, JA Jernigan, et al., "SHEA Guideline for Preventing Nosocomial Transmission of Multidrug-Resistant Strains of Staphylococcus aureus and Enterococcus," *Infection Control* and Hospital Epidemiology 24.5 (2003): 362-86.
- ³⁴ JM Boyce, "Do Infection Control Measures Work for Methicillin-Resistant Staphylococcus Aureus?" *Infection Control and Hospital Epidemiology* 25.5 (2004): 395-401.
- 35 SS Huang, et al., "Impact of routine intensive care unit surveillance cultures and resultant barrier precautions on hospital-wide methicillin-resistant Staphylococcus aureus bacteremia," *Clinical Infectious Diseases* 43.8 (2006): 971-8.
- ³⁶ Telephone interview with Dr. Carlene Muto, University of Pittsburgh Medical Center-Presbyterian Hospital, July 15, 2005 (412-629-2566).
- 37 "Skin Scourge: a Treatment-Resistant Form of Bacteria that Spreads Through Direct Contact, is Called a Greater Threat to Public Health than Sars or Bird Flu," Boston Globe, (August 21, 2006).

- ³⁸ Telephone interview with Dr. Carlene Muto, University of Pittsburgh Medical Center-Presbyterian Hospital, July 15, 2005 (412-629-2566).
- ³⁹ Based on an interview with Dr. Barry Farr by telephone, July 21, 2005.
- ⁴⁰ Dr. Richard Shannon, PowerPoint presentations, December 7, 2004 and March 11, 2005, Harvard Club of New York City; "PRHI in Progress: 2004 in Review," published by the Pittsburgh Regional Healthcare Initiative.
- ⁴¹ Dr. Richard Shannon, PowerPoint presentations, December 7, 2004 and March 11, 2005.
- 42 Ibid.
- 43 "Reducing hospital-acquired infections," Ivanhoe Broadcast News, (9-25-2004) at www. news8austin.com; "Simple Intervention Nearly Eliminates Catheter-related Bloodstream Infections," (12-09-2004) at www.sciencedaily.com
- 44 Joe Fahy, "Hospitals Able to Cut Some Kinds of Infections," Pittsburgh Post-Gazette (July 15, 2005) A1.
- ⁴⁵ Berenholtz, Sean M. et al, "Eliminating catheter-related infections in the intensive care unit." Critical Care Medicine Vol. 32, No. 10, Oct, 2004.
- 46 Ibid
- ⁴⁷ Pronovost, Peter et al. "An Intervention to Decrease Catheter-Related Bloodstream Infections in the ICU." *The New England Journal of Medicine*. Vol. 355, No. 26, December 26, 2006.
- ⁴⁸ E-mail communication between Dr. Brian Koll and Betsy McCaughey.
- ⁴⁹ Pronovost, Peter et al. "An Intervention to Decrease Catheter-Related Bloodstream Infections in the ICU." *The New England Journal of Medicine*. Vol. 355, No. 26, December 26, 2006.
- ⁵⁰ Berenholtz, Sean M. et al, "Eliminating catheter-related infections in the intensive care unit." Critical Care Medicine Vol. 32, No. 10, Oct, 2004.
- ⁵¹ E mail communication: B Koll to Betsymross@ aol.com 2-21-2008
- ⁵² Berenholtz, Sean M. et al, "Eliminating catheter-related infections in the intensive care unit." Critical Care Medicine Vol. 32, No. 10, Oct, 2004.
- ⁵³ "How Hospital Bug can Evade Attack", *BBC News*, http://news.bbc.co.uk June 25, 2006.
- McDonald, Clifford L. et. al, "Clostridium difficile Infection in Patients Discharged from US Short-stay Hospitals, 1996-2003. Emerging Infectious Diseases Vol. 12, No. 3, March, 2006.
- Obrien, Judith A., "The Emerging Infectious Challenge of Clostridium difficile Associated Disease in Massachusetts Hospitals", Infection Control and Hospital Epidemiology, Vol. 28, No. 11, November, 2007; Chandler, Rebecca E., "Clostridium difficile—Associated Disease in Oregon: Increasing Incidence and Hospital Level Risk Factors", Infection Control and Hospital Epidemiology, Vol. 28, No. 2, February 2007.
- Pèpin, Jaques et. al, "Mortality attributable to nosocomial Clostridium difficile-associated disease during an epidemic caused by a hypervirulent strain in Quebec.", Journal of the Canadian Medical Association, Vol. 173, No. 9, October 25, 2005; Warny, Michel et. Al, "Toxin production by an emerging strain of Clostridium difficile associated with outbreaks of severe disease in North America and Europe."
- Wenzel, Richard P., Prevention and Control of Nosocomial Infections, Lippincott Williams & Wilkins, 1997, pp.585-86; Bartlett, John G., "The New Clostridium difficile What Does It Mean?", The New England Journal Of Medicine, Vol. 353, No. 23, December 8, 2005
- 58 Walker, N. et al, "Blood pressure cuffs: friend or foe?", *Journal of Hospital Infection*, Vol. 63, Issue 2, June 2006, pp. 167-169.
- ⁵⁹ Muto, Carlene A. et al, "Control of an Outbreak of Infection with the Hypervirulent Clostridium difficile BI strain in a University Hospital Using a Comprehensive "Bundle" Approach", Clinical Infectious Diseases, Vol. 45 1266-73, 15 November, 2007.
- ⁶⁰ Flomenberg, Phyllis. Lecture at Thomas Jefferson University Medical Center Infection Control Conference, October 22, 2007.
- ⁶¹ Harper, LM, "Decreasing Clostridium difficile in the Newborn Intensive Care Unit Through Institution of Environmental Cleaning Procedures", Virox Solutions Newsletter, Vol. 14, 2007.
- 62 Ibid.
- ⁶³ Fakih, Mohamad G., et al., "Resident Physician (RP) and Physician Extender (PE) Survey Regarding Isolation Precautions for Patients with *Clostridium difficile*," Abstract 185 presented at SHEA's 17th Annual Scientific Meeting, April 2007.
- ⁶⁴ O'Brien, Judith A., et al., "The Emerging Challenge of Clostridium difficile-Associated Disease in Massachusetts Hospitals: Clinical and Economic Consequences." Infection

- Control and Hospital Epidemiology. Vol. 28, No.11, November 2007.
- 65 RP Shannon et al., "Economics of Central Line-Associated Bloodstream Infections," American Journal of Medical Quality 21.6 Supplement (2006): 78-16S.
- 66 Ibid.
- 67 Ibid.
- RR Roberts et al., "The Use of Economic Modeling to Determine the Hospital Costs Associated with Nosocomial Infections," *Clinical Infectious Diseases* 36.11 (2003) 1424-1432. This study puts the average cost of hospital infection at \$15,275; PW Stone et al., "A Systematic Audit of Economic Evidence Linking Nosocomial Infections and Infection Control Interventions, 1990-2000," *American Journal of Infection Control* 30.3 (2002): 145-52. This study estimates the average cost of infection to be \$13,973. Both studies look at incremental hospital costs only. The national cost was reached by multiplying 2 million infections by \$15,000 per infection.
- ⁶⁹ SE Cosgrove, Y Carmeli, "The Impact of Antimicrobial Resistance on Health and Economic Outcomes," *Clinical Infectious Diseases* 36.11 (2003): 1433-1437. The investigators found that antibiotic resistance increases length of stay, mortality rates, and costs.
- RR Roberts et al., "The Use of Economic Modeling to Determine the Hospital Costs Associated with Nosocomial Infections," *Clinical Infectious Diseases* 36.11 (2003) 1424-1432. The figure \$15,275 refers to excess hospital costs, not other patient care costs or economic losses incurred by the patient as a result of prolonged illness. The figure \$15,275 is similar to findings from other analyses of average hospital care costs due to infection. MedMined, Inc. analyzed the costs of infection using data from 50 hospital clients. It found that patients contracting infections in the hospital stayed an average of eight days extra and incurred \$14,000 in additional hospital costs. See: John Morrissey, "Debugging Hospitals," Modern Healthcare (April 26, 2004): 30.
- MM Peng et al., "Adverse Outcomes from Hospital-Acquired Infection in Pennsylvania Cannot be Attributed to Increased Risk on Admission," *American Journal of Medical Quality* 21.6 Supplement (2006): 17S-28S.
- Nancy M. Kane, Richard B. Siegrist, Jr., "Understanding Rising Hospital Inpatient Costs: Key Components of Cost and the Impact of Poor Quality," (August 12, 2002), unpublished manuscript. The findings are based on Medicare Cost Reports (1999-2000) and Uniform Hospital Discharge Data Set (2000) for all acute general hospitals in ten states: Massachusetts, New York, Virginia, Florida, Texas, Illinois, Iowa, California, Washington, and Colorado. Kane and Siegrist are adjunct faculty at the Harvard School of Public Health.
- 73 Ihid
- ⁷⁴ Institute for Healthcare Improvement, 100,000 Lives Campaign, How-to-Guide: Prevent Ventilator-Associated Pneumonia at www.lhi/org/IHI/Programs/Campaign.
- ⁷⁵ GA Noskin et al., "The Burden of *Staphylococcus aureus* Infections on Hospitals in the United States: an analysis of the 2000 to 2001 Nationwide Inpatient Sample Database," *Archives of Internal Medicine* 165.15 (2005): 1756-1761.
- ⁷⁶ Special thanks to Medmined Corporation (2006).
- 77 TE West, "Effect of targeted surveillance for control of methicillin-resistant Staphylococcus aureus in a community hospital system," *Infection Control and Hospital Epidemiology* 27.3 (2006): 233-8.
- ⁷⁸ H Grundmann et al., "Emergence and resurgence of meticillin-resistant Staphylococcus aureus as a public-health threat," *Lancet* 368,9538 (2006): 874-85.
- ⁷⁹ Gary A. Noskin, Robert J. Rubin, et.al., "National Trends in Staphylococcus aureus Infection Rates: Impact on Economic Burden and Mortality over a 6 Year Period (1998 to 2003), *Clinical Infectious Diseases* (2007) vol. 45: 1132-1140.
- 80 "Insurers Stop Paying for Care Linked to Error," Wall Street Journal (January 13, 2008)
- 81 Sanjay Saint, et.al., "Preventing Hospital-Acquired Urinary Tract Infection in the United States: A National Study," Clinical Infectious Diseases (January 15, 2008).
- ⁸² Demetriades, Demetrios et. Al., "When Does a Centers for Disease Control and Prevention Recommendation Become Standard of Care? Perhaps in the Courtroom. American College of Surgeons Mock Trial: Line Sepsis Liability" American College of Surgeons, published online 18 October, 2007.
- 83 Telephone interview with Dr. Frank Lowey, Professor of Microbiology, Columbia College of

- Physicians and Surgeons, July 15, 2005.
- 84 The CDC and the American Association of Medical Colleges are working on a Web-based curriculum that focuses largely on this issue.
- 85 Trick WE et al., "Impact of Ring Wearing on Hand Contamination and Comparison of Hand Hygiene Agents in a Hospital," *Clinical Infectious Diseases* 36.11 (2003): 1383-1390. Biant LC et al., "Eradication of *methicillin-resistant Staphylococcus aureus* by 'ring fencing' of elective orthopaedic beds," *British Medical Journal* 329.7458 (2004): 149-51. Visitors who sit on a chair or lean on a cabinet, then sit on the bed are transferring bacteria to the patient's bedclothes. CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," *Infections in Medicine* 20 (2003):194-200.
- Studies show that, nearly three quarters of patients' rooms are contaminated with MRSA and 69% with VRE. In one study, 42% of gloves worn by hospital personnel who had no direct patient contact but who touched contaminated surfaces became contaminated. JM Boyce et al., "Environmental contamination due to methicillin-resistant Staphylococcus aureus: possible infection control implications," Infection Control and Hospital Epidemiology 18.9 (1997): 622-627. A Concensus Statement by a multidisciplinary group of experts asked by the American Medical Association to provide guidelines for infection control cautions that: "In some cases caregivers actually go from patient to patient without changing their gloves, apparently confusing self-protection" with patient protection. DA Goldmann et al., "Strategies to Prevent and Control the Emergence and Spread of Antimicrobial-Resistant Microorganism in Hospitals," JAMA 275.3 (1996): 234-240.
- 87 Routine disinfection of stethoscopes between patients is recommended by the American Medical Association. CD Salgado, BM Farr, "MRSA and VRE: Preventing Patient-to-Patient Spread," *Infections in Medicine* 20 (2003):194-200; MA Marinella et al., "The stethoscope: a potential source of nosocomial infection?" *Archives of Internal Medicine*,157.7 (1997): 786-90; KC Zachary et al., "Contamination of gowns, gloves, and stethoscopes with vancomycin-resistant Enterococci," *Infection Control and Hospital Epidemiology* 22.9 (2001): 560-564; GA Noskin et al., "Recovery of vancomycin-resistant Enterococci on fingertips and environmental surfaces," *Infection Control and Hospital Epidemiology* 17.12 (1996): 770-772.
- The Agency for Healthcare Research and Quality recommends use of antibiotic catheters as one of its eleven patient safety practices. *Making Healthcare Safer: A Critical Analysis of Patient Safety Practices*. AHRQ Publication 01-E058, 2001. Also see: RO Darouiche et al., "A comparison of two antimicrobial-impregnated central venous catheters," *New England Journal of Medicine* 340.1 (1999): 1-8; I Raad et al., "Central venous catheters coated with Minocycline and Rifampin for the prevention of catheter-related colonization and blood-stream infections," *Annals of Internal Medicine* 127.4 (1997): 267-274.
- The following four studies support this suggestion: (1) MO Vernon et al., "Chlorhexidine gluconate to cleanse patients in a medical intensive care unit," *Archives of Internal Medicine* 166 (2006): 306-312. (2) LJ Hayek et al., "Preoperative whole body disinfection a controlled clinical study," *Journal of Hospital Infection* 11, Suppl. B (1988): 15-19 This study showed that two chlorhexidine showers reduced total infection rate by 30% and Staph aureus infections by 50%. (3) DJ Byrne et al., "Rationalizing whole body disinfection," *Journal of Hospital Infection* 15.2 (1990): 183-187. This study shows that a single shower does not maximize skin disinfection. The authors conclude that three showers should be recommended. (4) DS Paulson, "Efficacy Evaluation of a 4% Chlorhexidine Gluconate as a Full-Body Shower Wash," published by the Association for Practitioners in Infection Control (1993). This study found that showering for five days with chlorhexidine yielded maximum results for reducing bacteria on the skin, and keeping it low for 24 hours or more. "A 1 or 2 day presurgical application period is simply too short to establish the necessary levels of residual antimicrobial properties to be of value in reducing post-surgical infection rates."
- ⁹⁰ S Worcester, "Hospital system takes on MRSA," *Internal Medicine News* 38.19 (2005): 1-2.
- ⁹¹ A Kurz et al., "Perioperative Normothermia to Reduce the Incidence of Surgical-Wound Infection and Shorten Hospitalization," *New England Journal of Medicine* 334.19 (1996): 1209-1215.
- The Institute for Healthcare Improvement guidelines for improving infection prevention state that: "Administration of prophylactic antibiotics beginning 0 to 1 hour prior to surgical incision decreases the risk of surgical infection. http://www.ini.org/IHI/Topics/PatientSafety/ SurgicalSiteInfections/ImprovementStories (accessed 10-14-02). See also: JP Burke,

- "Maximizing appropriate antibiotic prophylaxis for surgical patients: an update from LDS Hospital, Salt Lake City," *Clinical Infectious Diseases* 33, Suppl. 2 (2001): S78-83.
- ⁹³ *Ibid.*, the Institute for Healthcare Improvement Guidelines for improving infection state that "surgical patients with core temperatures greater than 36 degrees C./ 98.6 degrees F are less likely to get an infection."
- 94 Ibid., the Institute for Healthcare Improvement states that "clipping instead of shaving results in decreased infection rates," and recommends that patients be told "not to shave the surgical site for 72 hours prior to surgery."
- 95 M A Ritter, "Operating Room Environment" Clinical Orthopaedics and Related Research 369 (1999): 103-109.
- ⁹⁶ Pittsburgh Regional Healthcare Initiative, "PHRI Executive Summary," (June, 2005).
- ⁹⁷ Urinary tract infections are the most common hospital-acquired infections. Limiting the use and duration of urinary tract catheters reduces risk of infection. See: J Puri et al., "Catheter Associated Urinary Tract Infections in Neurology and Neurosurgical Units," *Journal of Infection* 44.3 (2002): 171-175; F Stephan et al., "Reduction of Urinary tract infection and antibiotic use after surgery: a controlled, prospective, before-after intervention study," *Clinical Infectious Diseases* 24 (2006): 1544-1551.
- 98 CA Killian et al., "Risk Factors for Surgical-Site Infections Following Cesarean Section," Infection Control and Hospital Epidemiology 22.10 (2001): 613-7.
- 99 The CDC's National Nosocomial Infections Surveillance System, including the guarantees of confidentiality to hospitals, is described at www.cdc.gov/ncidod/hip/SURVEILL/NNIS.HTM
- ¹⁰⁰ Mark Chassin, "Achieving and Sustaining Improved Quality: Lessons from New York State and Cardiac Surgery," Health Affairs 21.4 (July/August 2002): 40-51.
- 101 *Ibid*.
- ¹⁰² JH Hibbard et al., "Does Publicizing Hospital Performance Stimulate Quality Improvement Efforts," *Health Affairs* 22.2 (March/April 2003): 84-94.
- ¹⁰³ For more information on the NNIS's methodology, consult: TC Horan & R. Gaynes, "Surveillance of nosocomial infections," *Hospital Epidemiology and Infection Control*, ed. CG. Mayhall (Philiadelphia: Lippincott, Williams & Wilkins, 2004) 1659-1702.
- ¹⁰⁴ J Steinhauer, "Hospitals in City Faulted for Failing to Report Many Errors," New York Times (February 13, 2001) B1. See also: M Chassin, EC Becher, "The Wrong Patient," Annals of Internal Medicine 136.11 (2002): 826-833.
- ¹⁰⁵ "Data Show Scourge of Hospital Infections," Washington Post, (July 13, 2005); "Hospitals Battling Infections," Washington Post, (July 14, 2005).
- 106 These are the types of infections that the Centers for Disease Control and Prevention recommend in a public reporting system. However, lawmakers could consider adding ventilator-related pneumonia, but these are more difficult to detect accurately, or urinary tract infections, which are common but less costly and deadly than other infections.
- ¹⁰⁷ O'Neill AJ, Cove JH, Chopra I. Mutation frequencies for resistance to fusidic acid and rifampin in Staphylococcus aureus. J Antimicrob Chemother 2001;47:647-650. *Unnecessary Deaths*, p.1.
- ¹⁰⁸ CA Muto, JA Jernigan, et al., "SHEA Guideline for Preventing Nosocomial Transmission of Multidrug-Resistant Strains of Staphylococcus aureus and Enterococcus," *Infection Control* and Hospital Epidemiology 24.5 (2003): 362-86.
- 109 Centers for Disease Control and Prevention. National Nosocomial Infection Surveillance (NNIS) System report: data summary from January 1992–June 2001. Am J Infect Control 2001;29:404-421.
- ¹¹⁰ Boyce JM, Potter-Bynoe G, Chenevert C, King T. Environmental contamination due to methicillin resistant *Staphylococcus aureus*: possible infection control implications. *Infect Control Hosp Epidemiol* 1997;18:622-627.
- ¹¹¹ Dr. Richard Shannon, PowerPoint presentations, December 7, 2004 and March 11, 2005.
- ¹¹² Galdbart JO, Morvan A, El Solh N. Phenotypic and molecular typing of nosocomial methicillin-resistant *Staphylococcus aureus* strains susceptible to gentamicin isolated from France from 1995-1997. *J Clin Microbiol* 2000; 8:185-190.
- ¹¹³ Bager F. DANMAP 98: Consumption of Antimicrobial Agents and Occurrence of Antimicrobials in Bacteria From Food Animals, Food and Humans in Denmark. Copenhagen: Statens Serum Institut, Danish Veterinary and Food Administration, Danish Medicines Agency, Danish Veterinary Laboratory; 1998.