ORIGINAL ARTICLE

Reconsidering Contact Precautions for Endemic Methicillin-Resistant Staphylococcus aureus and Vancomycin-Resistant Enterococcus

Daniel J. Morgan, MD, MS;¹ Rekha Murthy, MD;² L. Silvia Munoz-Price, MD, PhD;³ Marsha Barnden, RNC, MSN, CIC;⁴ Bernard C. Camins, MD, MSc;⁵ B. Lynn Johnston, MD, MSc;⁶ Zachary Rubin, MD;⁷ Kaede V. Sullivan, MD;⁸ Andi L. Shane, MD, MPH, MSc;⁹ E. Patchen Dellinger, MD;¹⁰ Mark E. Rupp, MD;¹¹ Gonzalo Bearman, MD, MPH¹²

BACKGROUND. Whether contact precautions (CP) are required to control the endemic transmission of methicillin-resistant *Staphylococcus aureus* (MRSA) or vancomycin-resistant *Enterococcus* (VRE) in acute care hospitals is controversial in light of improvements in hand hygiene, MRSA decolonization, environmental cleaning and disinfection, fomite elimination, and chlorhexidine bathing.

OBJECTIVE. To provide a framework for decision making around use of CP for endemic MRSA and VRE based on a summary of evidence related to use of CP, including impact on patients and patient care processes, and current practices in use of CP for MRSA and VRE in US hospitals.

DESIGN. A literature review, a survey of Society for Healthcare Epidemiology of America Research Network members on use of CP, and a detailed examination of the experience of a convenience sample of hospitals not using CP for MRSA or VRE.

PARTICIPANTS. Hospital epidemiologists and infection prevention experts.

RESULTS. No high quality data support or reject use of CP for endemic MRSA or VRE. Our survey found more than 90% of responding hospitals currently use CP for MRSA and VRE, but approximately 60% are interested in using CP in a different manner. More than 30 US hospitals do not use CP for control of endemic MRSA or VRE.

CONCLUSIONS. Higher quality research on the benefits and harms of CP in the control of endemic MRSA and VRE is needed. Until more definitive data are available, the use of CP for endemic MRSA or VRE in acute care hospitals should be guided by local needs and resources.

Infect. Control Hosp. Epidemiol. 2015;36(10):1163-1172

Despite decades of experience, the use of contact precautions (CP) for endemic methicillin-resistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant *Enterococcus* (VRE) remains controversial.^{1,2} As a result, there is a growing diversity of practice for CP in acute care hospitals.^{1,3} A North American group of adult and pediatric hospital epidemiologists and infection prevention experts with expertise in guideline development met on the Society for Healthcare Epidemiology of America (SHEA) Guidelines committee and, independent of SHEA or SHEA endorsement, completed this article to elucidate the current state of the literature pertaining to the application and discontinuation of CP for endemic MRSA and VRE. In addition, the group administered a survey to the SHEA Research Network of hospital epidemiologists and infection preventionists to better ascertain the practice and

experience with CP for endemic MRSA and VRE. Finally, a convenience sample of hospitals that do not use CP for MRSA or VRE was identified from the literature and an infection control listserv, and their experiences were elicited and summarized.

METHODS

Guidelines were reviewed for recommendations relating to use of CP for endemic MRSA or VRE. A literature search for English language publications from 2003 through 2013 was conducted on PubMed using the search terms "CP," "barrier precautions," "isolation," "MRSA," and "VRE" to identify papers that compared the use of CP with some other standard for the control of MRSA and VRE in endemic settings.

Affiliations: 1. University of Maryland, Baltimore, Maryland; 2. Cedars-Sinai Medical Center, Los Angeles, California; 3. Medical College of Wisconsin, Milwaukee, Wisconsin; 4. Adventist Health System, Roseville, California; 5. University of Alabama at Birmingham, Birmingham, Alabama; 6. Dalhousie University, Halifax, Nova Scotia; 7. David Geffen School of Medicine at UCLA, Los Angeles, California; 8. Clinical Microbiology Laboratory, Children's Hospital of Philadelphia, Philadelphia, Pennsylvania; 9. Emory University School of Medicine, Atlanta, Georgia; 10. Department of Surgery, University of Washington Medical Center, Seattle, Washington; 11. University of Nebraska Medical Center, Omaha, Nebraska; 12. Medical College of Virginia, Richmond, Virginia.

Received February 24, 2015; accepted June 5, 2015; electronically published July 3, 2015

^{© 2015} by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2015/3610-0006. DOI: 10.1017/ice.2015.156

Publications focusing on outbreak settings were excluded. A survey was mailed electronically to all SHEA Research Network members. Hospitals not using CP for MRSA or VRE were identified from both the literature and an infection control listserv and queried on practice and experience; we summarize only reports previously published or with permission from the institutions.

RESULTS

Guideline Recommendations for CP for MRSA and VRE in Acute Care Facilities

Multiple guidelines address strategies for preventing crosstransmission of MRSA and VRE in acute care settings that reference the use of CP. SHEA and the Infectious Diseases Society of America jointly recommend that CP be used for MRSA-infected and MRSA-colonized patients in acute care settings for the control of MRSA in both endemic and outbreak settings.⁴ More broadly, the Healthcare Infection Control Practices Advisory Committee and the Centers for Disease Control and Prevention recommend that CP be implemented routinely in "all patients infected with target MDROs [multidrug-resistant organisms] and for patients that have been previously identified as being colonized with target MDROs" without identifying explicitly which MDROs are to be included.⁵

Impact of CP on Endemic MRSA

Forty-eight articles were reviewed by 2 individuals (Z.R. and B.C.C.) and final results discussed by all authors regarding MRSA. CP as an intervention to decrease MRSA acquisition was rarely analyzed separately from other interventions, and most studies were performed in outbreak settings where multiple control measures were initiated simultaneously. Initially, only studies that evaluated CP alone were included in the review. However, only 2 studies in endemic settings qualified for inclusion, one of which was a prospective quasi-experimental study^{6,7} and one a randomized trial.⁸ Given the paucity of studies evaluating the effect of CP alone, we then included other studies that evaluated the effect of active surveillance cultures (ASC) and resultant increase in use of CP^{9–15} or universal gown and gloves.¹⁶

Lower quality, quasi-experimental studies generally demonstrated a decrease in transmission of MRSA with CP. In a retrospective analysis of interventions to decrease MRSA bacteremia, authors concluded that CP and ASC resulted in a 67% decrease in the incidence of MRSA bacteremia.¹³ MRSA acquisition decreased from 7.0% to 2.8% after implementation of similar interventions in another quasi-experimental study.¹⁴ In a larger quasi-experimental study, Robicsek et al¹⁵ instituted ASC and CP on all hospital admissions with a subsequent decrease in MRSA. This study included a decolonization regimen in its final phase. Marshall and colleagues¹⁰ performed a quasi-experimental study in an intensive care unit (ICU) with endemic MRSA and noted decreased rates of MRSA after changing from no-CP to CP-based ASC. Another before-after study compared 4 different infection prevention strategies and demonstrated a decrease in MRSA bacteremia with CP.¹³ Finally, all hospitals of the US Department of Veterans Affairs implemented a before-after bundle that included CP based on ASC, hand hygiene, and cultural change. This study found a small decrease in MRSA colonization and a larger decrease in MRSA healthcare-associated infections.¹⁷

In contrast to uncontrolled studies, prospective trials with control groups largely failed to demonstrate a benefit of CP for MRSA. In a prominent controlled quasi-experimental study, Harbarth et al9 screened surgical patients for MRSA colonization at admission. Using a cross-over design in 12 surgical wards, they compared rapid ASC with CP to standard infection control measures, which included less frequent CP and decolonization for patients with MRSA by clinical cultures. They observed no difference in MRSA rates between the 2 periods (adjusted incidence rate ratio, 1.20 [95% CI, 0.85–1.69]; P = .29). Huskins et al¹² conducted a multicenter cluster randomized controlled trial examining ASC and CP for MRSA-colonized patients and found no difference in the incidence of MRSA colonization or infection. A 2014 study conducted across 13 European ICUs evaluated multiple interventions for MDROs in a quasi-experimental fashion. The final phase of the study evaluated ASC with application of CP for carriers.¹¹ The authors found that colonization with MDROs (MRSA, VRE, and Enterobacteriaceae) decreased slightly during an earlier chlorhexidine and hand hygiene intervention phase of the study (relative risk, 0.98 [95% CI, 0.95-0.99]; P=.04) but did not decrease with subsequent addition of ASC.⁶

Studies examining the use of universal gloves or universal gowns and gloves have identified mixed results, with the largest study identifying a decrease in MRSA transmission.¹⁶ However, in units randomized to universal gowns and gloves, the number of patient interactions by healthcare personnel (HCP) was lower with better hand hygiene and thus the decreased transmission of MRSA may have been due only indirectly to gown and glove use.¹⁶ In a quasi-experimental study comparing CP for MRSA versus universal gloving, Bearman et al⁶ showed no difference in MRSA acquisition. Harris and colleagues¹⁶ published a cluster randomized trial in which the use of universal gloves, regardless of colonization status, decreased MRSA acquisition by 40%.

In summary, many studies suffer from methodologic limitations, such as small sample size, interventions introduced simultaneously, and lack of comparison groups. Adherence to CP was often not monitored, and when assessed, adherence was poor (Table 1a). Although retrospective studies suggest that CP decreases MRSA acquisition, this was not observed in more rigorous studies.

Impact of CP on Endemic VRE

Forty-five articles were reviewed by 2 individuals (M.B. and B.L.J.) for VRE. The literature $^{18-33}$ abounds with publications

TABLE 1A.	Literature Review of Articles Fron	n 2004 to 2013 That Examined the Effect of CI	P (With or Without Other Measures) on MRSA
-----------	------------------------------------	---	--

			Interventions used						
Lead author	Trial design	Setting	Gowns	Gloves	Surveillance Culturing	НН	Universal decolonization	Targeted decolonization	Main findings
Trick et al ⁸	RCT	SNFs			-	-	_	_	UG use was equivalent to CP in SNFs that did not limit patient activities
Lucet et al ¹⁴	Before-after	ICUs				-	-	-	Surveillance cultures to guide CP led to a decrease in MRSA acquisition rates
Huang et al ¹³	Quasi- experimental	ICUs	\checkmark	\checkmark	\checkmark	-	_	_	Surveillance cultures to guide CP decreased MRSA acquisition rates and BSI rates; same decrease in BSI rates observed hospital-wide
Robicsek et al ¹⁵	Before-after	Hospital-wide	\checkmark	\checkmark	\checkmark	-	-	\checkmark	Surveillance cultures to guide CP and targeted colonization resulted in a decrease in invasive MRSA infection rates
Harbarth et al ⁹	Cross-over quasi- experimental	Surgical patients	\checkmark	\checkmark	\checkmark	-	-	\checkmark	Surveillance cultures to guide CP and targeted decolonization did not reduce nosocomial MRSA infection rates with endemic MRSA prevalence
Bearman et al ³⁴	Before-after	ICUs	-		-		-	-	UG use was equivalent to CP for prevention of MRSA acquisition
Huskins et al ¹²	RCT	ICUs	\checkmark		\checkmark	-	-	-	Surveillance cultures to guide CP vs standard CP alone resulted in equivalent MRSA acquisition or infection rates
Jain et al ¹⁷	Before-after	Hospital-wide	\checkmark	\checkmark	\checkmark		_	_	Bundle of surveillance cultures to guide CP, HH, and institutional culture change was associated with a decrease in MRSA colonization and infection rates
Derde et al ⁶⁸	RCT	ICUs							No impact of surveillance cultures to guide CP
Harris et al ¹⁶	RCT	ICUs	v	v	v	-	• _	<u> </u>	Universal CP use significantly reduced MRSA acquisition
Marshall et al ¹⁰	Before-after	ICUs			$\dot{\checkmark}$	-	-	-	Surveillance cultures to guide CP resulted in a decrease in MRSA acquisition rates

NOTE. BSI, bloodstream infection; CP, contact precautions; HH, hand hygiene; ICU, intensive care unit; MRSA, methicillin-resistant *Staphylococcus aureus*; RCT, randomized controlled trial; SNF, skilled nursing facility; UG, universal gloving.

	Trial design	Setting	Interventions used						
Lead author			Gowns	Gloves	Surveillance cultures	HH	Universal decolonization	Targeted decolonization	Main findings
Bearman et al ⁶	Before-after	MICU	Before		\checkmark		No	No	No difference in VRE acquisition risk between CP and UG use
Bearman et al ³⁴	Before-after	SICU	Before	\checkmark	\checkmark	\checkmark	No	No	No difference in VRE acquisition risk between CP and UG use
Huskins et al ¹²	RCT of 18 ICUs	ICU	\checkmark	\checkmark	\checkmark	\checkmark	No	No	No impact of surveillance culturing and isolation for MDROs
Harris et al ¹⁶	RCT of 20 ICUs	ICUs	\checkmark	\checkmark	_	-	-	-	Universal CP use had no effect on VRE acquisition but was associated with less MRSA acquisition
Derde et al ¹¹	Before-after	ICU	\checkmark	\checkmark	\checkmark	\checkmark	\checkmark	No	No impact of surveillance culturing and isolation for MDROs

TABLE 1B. Literature Review of Articles From 2004 to 2013 That Examined the Effect of CP (With or Without Other Measures) on VRE

NOTE. CP, contact precautions; HH, hand hygiene; ICU, intensive care unit; MDRO, multidrug-resistant organism; MICU, medical intensive care unit; MRSA, methicillin-resistant *Staphylococcus aureus*; RCT, randomized controlled trial; SICU, surgical intensive care unit; UG, universal gloving; VRE, vancomycin-resistant *Enterococcus*.

reporting the benefit of CP in terminating VRE outbreaks. As with the MRSA literature, CP as an intervention to decrease VRE acquisition was rarely studied separately from other interventions or compared with standard precautions as the only intervention. Therefore, reviewers included studies that compared CP alone or with some other intervention with a defined control. The search for published studies examining use of CP for VRE control in non-outbreak settings identified 5 studies (Table 1b).

Bearman et al^{6,34} conducted 2 quasi-experimental studies where CP for patients with VRE was compared with universal glove use. The authors found no difference in VRE acquisition and higher healthcare-associated infection rates with universal glove use in one of the studies. In 2014, De Angelis et al³⁵ published a systematic review and meta-analysis of measures taken to control VRE in ICU settings. They reported results from 3 studies^{6,12,36} that had application of CP as their only intervention. CP did not significantly reduce the VRE acquisition rate (pooled relative risk, 1.08 [95% CI, 0.63–1.83]).

The remaining 3 studies were cluster-randomized trials that examined the impact of CP on VRE acquisition in ICUs.^{11,12,16} Huskins et al¹² used CP in the intervention group after ASC. The mean ICU-level incidence of colonization or infection with VRE/1,000 patient-days at risk did not differ between the 2 groups (P=.53). In a cluster randomized trial among ICUs, HCP in intervention ICUs wore gowns and gloves for all patient contacts and room entries in comparison with control ICUs where CP was used only for patients with known antibioticresistant bacteria, and the researchers found no difference.¹⁶ Likewise, a study in the setting of universal chlorhexidine body washes and hand hygiene improvement identified no benefit to ASC for addressing VRE or other MDROs.¹¹

In conclusion, the literature has not identified a benefit to CP over standard precautions in acute care settings for controlling the spread of VRE. Unfortunately, no study has compared CP with standard precautions alone. Positive publication bias likely exists and study quality is generally low.

Studies in Children

Studies assessing the impact of CP for MRSA or VRE in children are limited to quasi-experimental studies in outbreak settings.²⁰ A case-control study with 16 cases and 62 controls identified the absence of CP (odds ratio, 17.16 [95% CI, 1.49–198.21]) and the presence of a gastrointestinal device (4.03 [1.04–15.56]) as factors associated with VRE acquisition.²⁵ As with adult studies, the pediatric literature is limited to quasi-experimental studies that examined CP as part of a bundle, often in response to an outbreak.²⁰

Potential Harms Associated With CP for MRSA and VRE

Studies exploring the negative consequences of CP have focused on the impact on HCP behavior, patient flow, adverse physical events, psychological harm, and patient satisfaction. Various studies have examined the impact of CP on HCP behavior.^{3,37–41} CP has been associated with fewer bedside visits and physical examinations by HCP. In ICU and medical/ surgical wards at 4 hospitals, patients on CP were observed having fewer hourly HCP visits (2.78 vs 4.37; P < .001) and shorter contact time (14.0 vs 17.0 minutes/hour; P = .02).³ In surgical settings, patients on CP received 5.3 hourly visits compared with 10.9 among patients not on CP, and had a shorter contact time (29 vs 37 minutes/hour; P = .008).³⁷ In a medical ICU, patients on CP had fewer contacts than those who were not on CP (2.1 vs 4.2 per hour; P = .03).³⁸ Similarly, attending physicians examined patients on CP less frequently (35% vs 73%; P < .001).⁴¹

Studies suggest that CP may delay admission from emergency to inpatient settings. Duration of time for admission from the emergency department to a CP room was 12.9 hours for patients with MRSA compared with 10.4 hours for a standard room.⁴² Average admission wait was 54 minutes longer in patients with a history of MDRO (298 minutes vs 244 minutes; P = .045).⁴³ CP may also result in delayed discharge of patients. Patients on CP awaiting transfer to longterm care facilities experienced an average delay of 10.9 days compared with 4.3 days for similar patients not on CP.^{42–44}

A retrospective study at 2 tertiary medical centers found adverse event rates were higher in patients on CP (31/1,000 patient-days vs 15/1,000 patient-days; P < .001) as were preventable adverse event rates (20/1,000 patient-days vs 3/1,000 patient-days; P < .001).⁴⁵ Karki et al⁴⁶ studied inpatients before and after application of CP for positive VRE status and found no difference in rates of adverse events (incidence rate ratio, 1.04 [0.85-1.27]) but sub-analyses noted more injuries after CP were initiated (3.24 [1.16–11.17]).⁴⁶ By contrast, in a case-control study, patients with MRSA (on CP) with heart failure or chronic obstructive pulmonary disease found no difference in complication rates with patients without MRSA (P = .40).⁴⁰ Notably, 2 trials that randomly applied CP to patients regardless of MDRO status found no increase in adverse events associated with use of CP.^{16,39} Additionally, 2 studies failed to find differences in morbidity or complications in patients on CP and those that were not.^{39,40}

There is a significant quantity of literature related to psychological and psychiatric outcomes in patients on CP but findings vary.^{47–56} Among inpatients at 3 general hospitals, patients on CP had higher Hospital Anxiety and Depression Scale scores (12.8 vs 8.2; P < .001).⁵¹ For patients on a spinal cord injury rehabilitation unit, those on CP had higher Beck Depression Inventory scale scores (16.5 vs 12.3; P = NS).⁵² Subsequent controlled studies by Day et al^{53,56} suggest that CP may not be associated with depression and anxiety.

The issue of isolation is relevant to the care of pediatric patients, who may be unable to visit unit playrooms or schoolrooms in hospitals owing to their isolation status. Despite these potential concerns, one study in pediatrics found no difference in care.⁵⁷

Although a number of studies have investigated the relationship between CP and patient satisfaction, patient perceptions about the quality of care varied.^{58,59} In medical and surgical inpatient wards, Mehotra et al⁵⁸ found that patients on CP were more likely to have concerns with their care than patients who were not on CP (odds ratio, 2.0 [95% CI, 1.3–3.2]). In contrast, Gasink et al⁵⁹ administered the Consumer Assessment of Healthcare Providers and Systems Hospital Survey to medical and surgical inpatients exposed and unexposed to CP and reported that CP was not associated with less satisfaction.

In conclusion, CP consistently appears to modify HCP behavior, leading to fewer patient contacts. Multiple types of harm have been described with CP in the literature but results have been inconsistent and study quality has been relatively low.

Proportion of Patients on CP for MRSA or VRE

CP is applied to a substantial proportion of hospitalized patients and varies by geographical area and the methods used to identify MRSA or VRE. If samples obtained during routine clinical care are the basis for identifying MRSA or VRE, an estimated 5%–10% of patients in US acute-care facilities are isolated compared with 20%–25% of patients when surveillance testing for MRSA or VRE is used to identify colonization.^{12,16,17,60–62} Because patients on CP have longer lengths of

hospital stay, the proportion of patients on CP on a ward can be as high as 60%.⁶¹

Survey of SHEA Members/SHEA Research Network on Use of CP

The SHEA Research Network is an international consortium of more than 200 hospitals conducting multicenter research projects in healthcare epidemiology. A total of 87 members of the SHEA Research Network responded to the survey regarding their institutions' use of CP for MRSA and VRE (response rate, 33% [87/267]). Table 2 summarizes respondent perceptions and attitudes toward CP. Most respondents worked at acute care hospitals (93%) and belonged to teaching or teaching-affiliated hospitals (72%). Ninety-two percent of respondents reported using CP in their respective facilities for both MRSA and VRE. Respondents applied CP for positive surveillance screens (nasal, axillary, or perineal screen) for MRSA (48%) and VRE (49%), diarrhea (71%), uncontrolled secretions (44%), and uncovered wounds (27%). Cohorting of MRSA- or VRE-colonized patients in double occupancy rooms was either never done (46%) or performed only in extreme cases of bed shortage (43%). Most respondents (63%) were in favor of implementing CP in a different fashion than current practice (Figure 1), and most felt that CP decreased the number of HCP visits to patients (78%) and had a negative impact on mental

TABLE 2. Results of SHEA Research Network Survey of Respondents' Beliefs Relating to CP

Extent to which HCP believe CP prevents			
	Have a large impact	Have a slight impact	Have no Impac
MRSA	31 (41%)	36 (47%)	9 (12%)
VRE	27 (36%)	38 (51%)	9 (14%)
Ways in which HCP believe CP causes harm			
Decrease in number of visits	58 (78%)		
Negative impact on patient's mental health	46 (68%)		
Negative impact on patient's satisfaction	50 (69%)		
Increase in adverse events (eg, falls or pressure ulcers) ^a	26 (38%)		
HCP opinion of CP	Dislike	Like	
Physicians	61 (94%)	3 (5%)	
Nurses	48 (76%)	9 (4%)	
Others	52 (87%)	4 (7%)	
Beliefs regarding routine use of surveillance culturing and CP for MRSA and VI	RE		
Routine surveillance culturing and CP helpful in ICUs	24%		
Targeted surveillance culturing and CP helpful in wards or high-risk population	18%		
Surveillance culturing and CP useful during outbreaks	32%		
Surveillance culturing and CP not helpful	21%		

NOTE. CP, contact precautions; HCP, healthcare personnel; ICU, intensive care unit; MRSA, methicillin-resistant *Staphylococcus aureus*; SHEA, Society for Healthcare Epidemiology of America; VRE, vancomycin-resistant *Enterococcus*.

^aAdditional responses: decrease 16 (24%), no impact 26 (38%).

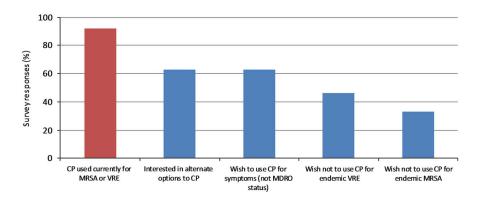


FIGURE 1. Results from Society for Healthcare Epidemiology of America Research Network survey respondents regarding opinions for use of contact precautions (CP). MDRO, multidrug-resistant organisms; MRSA, methicillin-resistant *Staphylococcus aureus*; VRE, vancomycin-resistant *Enterococcus*.

TABLE 3. Practices Being Used in Place of Standard Centers for Disease Control and Prevention Contact Precautions for Patients Identified With MRSA or VRE by a Convenience Sample of Hospitals in the United States

Institution (number of hospitals)	MRSA	VRE	C. difficile	MDR-GNR	Year foregoing CP
Hospitals that practice enhanced focus on hand hygi	ene complianc	e and HAI	prevention bund	les (horizontal inte	erventions)
Virginia Commonwealth University MC	No	No	Yes	Yes	2013
University of Massachusetts (2 hospital campuses)	No	No	Yes	Yes	2010
Detroit MC (7 hospitals)	No	No	Yes	Yes	Prior to 2003
Tufts-New England MC	No	No	Yes	Yes	2010
St. Johns MC, Santa Monica, CA	No	No	Yes	Yes	2002
University of Rochester MC	No	No	Yes	Yes	2014
Baylor St. Luke's MC	No ^a	No	Yes	Yes	2005
UCLA (2 hospitals)	No	No	Yes	Yes	2013
University of Nebraska MC	No	No	Yes	Yes	2015
San Francisco General Hospital	No	No	Yes	Yes	Prior to 2002
University of San Francisco MC	No	No	Yes	Yes	Prior to 2002
Alta Bates MC, Oakland, CA	No	Yes	Yes	Yes	2014
University of Cincinnati MC	No	Yes	Yes	Yes	Prior to 2002
Oakwood Hospital System, MI (4 hospitals)	No	No	Yes	Yes	Prior to 2013
Hospitals that use gowns and gloves for syndromic i	ndications onl	y (diarrhea,	draining wound	s)	
Baystate Hospitals (multiple hospitals) ²	No	No	Yes ^b	Yes	2003
Dartmouth MC ²	No	No	Yes ^b	Yes	Prior to 2003
Hospitals that use decolonization of patients identifi	ed to have S. a	ureus (inclu	iding MRSA) ^c		
Cleveland Clinic (10 hospitals)	No	No	Yes	Yes	Prior to 2003

NOTE. All institutions agreed to publication of their name and practice or were in the published literature. *C. difficile, Clostridium difficile*; CP, contact precautions; HAI, healthcare-associated infection; HCP, healthcare personnel; MA, Massachusetts; MC, Medical Center; MDR-GNR, multidrug-resistant gram-negative rods including carbapenem-resistant *Acinetobacter* and carbapenem-resistant Enterobacteriaceae; MRSA, methicillin-resistant *Staphylococcus aureus*; UCLA, University of California, Los Angeles; VRE, vancomycin-resistant *Enterococcus*. ^aOnly if MRSA present in wounds with uncontained drainage are patients placed on contact precautions.

^bUse gloves without gowns for all patients with diarrhea regardless of *C. difficile* testing.

^cDecolonization consisted of chlorhexidine bathing and intranasal mupirocin.

health (68%) and on patient satisfaction (69%). In addition, a high proportion of respondents (63%) were in favor of employing CP for symptoms such as diarrhea, draining wounds, and uncontrolled secretions, regardless of MDRO status.

Alternative Approaches to CP for Endemic MRSA or VRE

Most US hospitals use CP for endemic MRSA or VRE. However, some hospitals are not using CP for MRSA or VRE but are employing different approaches. Approaches to MRSA or VRE control that do not use CP generally fall into 3 categories: (1) focus on improved general or horizontal infection control methods without CP, (2) enhanced efforts on syndromic use of gowns and gloves for patients with syndromes correlated with greater contamination (eg, diarrhea, wounds), and (3) targeted decolonization of patients found to be positive for MRSA without CP (see Table 3).

Several institutions (Table 3) focus on general horizontal approaches to limiting transmission of MRSA and VRE, such as hand hygiene, bathing patients with chlorhexidine, or environmental cleaning and disinfection. These hospitals continue to apply CP for *Clostridium difficile* and multidrug-resistant gram-negative rods. There were multiple anecdotal reports from these institutions of stable or declining rates of infections with MRSA or VRE after foregoing CP.^{63–65}

Three centers reported using CP for patients with specific syndromes regardless of colonization status. These centers made a specific effort to use CP for all patients with diarrhea who were unable to self-toilet or with incontinence (including *C. difficile* or norovirus), open wounds that cannot be contained within a dressing, pneumonia or upper respiratory tract infection in patients unable to practice respiratory etiquette, and patients with urinary tract infection unable to self-toilet⁶⁵ or with incontinence.⁶⁶ The limited reports from these hospitals noted no change in percentage of *S. aureus* that is methicillin-resistant, a low rate of MRSA during a prevalence survey, and stable or declining rates of ventilator-associated pneumonia, central line–associated bloodstream infection, and surgical site infection (both overall and due to MRSA).⁶⁵

Given the importance of preventing infections with either methicillin-susceptible and methicillin-resistant *S. aureus*, the Cleveland Clinic hospital system implemented surveillance cultures of patients for *S. aureus* upon admission to ICU with targeted decolonization with chlorhexidine bathing and intranasal mupirocin. They reported decreased *S. aureus* in a single medical ICU (6.28 vs 3.32 acquisitions/1,000 patient-days) and healthcare-associated infections (3.52 to 1.29 cases/1,000 patient-days).⁶⁷ This policy has since been implemented at all 10 Cleveland Clinic hospitals with reported declining rates of MRSA. These hospitals continue to apply CP for *C. difficile* and multidrug-resistant gram-negative rods.

DISCUSSION

The literature does not provide strong evidence of benefit from CP over standard precautions for controlling endemic VRE or MRSA. To our knowledge, to date, no study has compared CP with standard precautions. Determining the optimal use of CP is an important issue because it affects 10%–25% of hospitalized patients, may have a negative impact on patient throughput, and may cause harm and decrease quality of care by reducing HCP-patient contact. Understanding the true benefits and harms of CP is important. Our survey of SHEA Research Network members found that most hospitals responding currently use CP for MRSA and VRE, but a high

proportion expressed interest in using CP in a different manner.

Hospitals no longer using CP for MRSA or VRE paid special attention to collecting metrics focusing on processes and outcomes. Process measures generally focused on HCP compliance with policies related to hand hygiene and use of gloves and gowns, as well as compliance with other horizontal infection control strategies being employed at each institution (eg, hand hygiene improvement, line insertion checklists, chlorhexidine bathing, environmental cleaning, and antimicrobial stewardship). In addition, the availability of single patient rooms was reported by some facilities to factor in the decision to not routinely use CP for MRSA and VRE. Outcome measures focused on overall, hospital-wide rates of healthcareassociated infections, especially those due to MRSA or VRE. A few facilities conducted either limited or ongoing surveillance culturing for MRSA patient colonization to ensure that MRSA and VRE rates did not increase after foregoing CP.

Surprisingly, hospitals not using CP for patients with MRSA or VRE reported no negative feedback from the Joint Commission or the Centers for Medicare and Medicaid Services after hospital visits. Because not using CP for MRSA or VRE is uncommon, many respondents stated that they had been proactive in providing data to surveyors related to MRSA and VRE rates and having infection prevention policies that clearly stated the rationale for not using CP for MRSA or VRE. At all institutions it was important that staff be educated with regard to use of gowns and gloves so that they would be compliant with policies and could explain policies if asked by regulatory reviewers. Interestingly, some hospitals designed their program to forego CP with assistance from the local and state Departments of Health and reported that this step assisted with regulatory review.

It is notable that many hospitals not using CP for MRSA or VRE are in states with legislation mandating active surveillance culturing for MRSA. Despite mandating use of active surveillance, state laws often do not require use of CP for those identified with MRSA. This was not seen as a barrier to foregoing use of CP.

Relevant questions for future research include when and where CP may provide additional benefits over assiduous use of standard precautions, especially when hospitals are using horizontal control measures, such as chlorhexidine bathing, universal gloving, hand hygiene surveillance, and environmental cleaning. Additionally, a more rigorous examination of universal or targeted chlorhexidine bathing or syndromic use of CP compared with standard use of CP for MRSA or VRE would advance the field. Our findings suggest that a "one size fits all" approach to MRSA and VRE control in endemic settings is not supported by robust science. Across multiple healthcare systems, various strategies are reported for the control of endemic, hospital-acquired MRSA and VRE infections, suggesting that local factors, needs, and resources should drive the choice of optimal CP utilization.

In conclusion, no high quality data support the use of CP for endemic MRSA or VRE and there may be patient harms and unintended consequences associated with CP. The burden of CP is not insignificant because approximately 25% of hospitalized patients are on CP when surveillance culturing is employed. Although most US hospitals currently use CP for MRSA and VRE, a high proportion of SHEA Research Network respondents expressed interest in foregoing CP for the control of endemic MRSA and VRE. At least 30 US hospitals do not use CP for endemic MRSA or VRE and generally rely on broad-based, bundled interventions such as hand hygiene, chlorhexidine bathing, environmental cleaning, and checklists. Higher quality research on the risks and benefits of CP is needed. Until more definitive data are available, the use of CP for control of endemic MRSA or VRE in acute care hospitals should be guided by local needs and resources.

ACKNOWLEDGMENTS

We thank Valerie Deloney for her invaluable assistance with the preparation of this article.

Financial support. SHEA Research Network.

Potential conflicts of interest. D.J.M. reports that he has an advisory/consultant role for Welch Allyn, Sanogiene/Biomed, and 3M. B.C.C. reports that he has received research grants/contracts from Pfizer, Merck, and the Centers for Disease Control and Prevention (Preventing Hemodialysis-Related Bloodstream Infections). E.P.D. reports that he has an advisory/consultant role and has received honoraria from Merck, Baxter, Ortho-McNeil, Targanta, Schering-Plough, Astellas, CareFusion, Durata, Pfizer, and Rib-X; and research grants/contracts from Exoxemis. B.L.J. reports that she has received research grants/contracts from Gilead Sciences and Pfizer Canada, education grants from GlaxoSmithKline, Sunovion Pharmaceuticals Canada, Optimer Pharmaceuticals Canada for Infectious Diseases Continuing Medical Education for Family Physicians (October 2013), and grants from Merck Canada, ViiV Healthcare Canada, Bristol-Myers Squibb Canada, and Gilead Sciences for Atlantic Canada Human Immunodeficiency Virus Educational Conference. R.M. reports that she has received research grants/contracts from Medimmune. K.V.S. reports that he has received research grants/contracts from Nanosphere, Techlab, and Premier. G.B. reports that he has received research grants from Pfizer, Cardinal Healthcare, BioVigil, and Vestagen. M.E.R. reports that he has received research grants/contracts from National Institutes of Health, 3M, Magnolia; Consultant, and Sharklet. All other authors report no conflicts of interest relevant to this article.

Address correspondence to Daniel J. Morgan, MD, MS, 685 W. Baltimore St, MSTF 334, University of Maryland, Baltimore, MD 21201 (dmorgan@epi. umaryland.edu).

REFERENCES

- 1. Fätkenheuer G, Hirschel B, Harbarth S. Screening and isolation to control meticillin-resistant *Staphylococcus aureus*: sense, nonsense, and evidence. *Lancet* 2014;385:1146–1149.
- Morgan DJ, Kaye KS, Diekema DJ. Reconsidering isolation precautions for endemic methicillin-resistant *Staphylococcus aureus* and vancomycin-resistant. *Enterococcus. JAMA* 2015;312: 1395–1396.
- Morgan DJ, Pineles L, Shardell M, et al. The effect of contact precautions on healthcare worker activity in acute care hospitals. *Infect Control Hosp Epidemiol* 2013;34:69–73.

- Calfee DP, Salgado CD, Milstone AM, et al. Strategies to prevent methicillin-resistant *Staphylococcus aureus* transmission and infection in acute care hospitals: 2014 update. *Infect Control Hosp Epidemiol* 2014;35:772–796.
- CDC/HICPAC. Prevention of transmission of multidrug resistant organisms 2009. www.cdc.gov/hicpac/mdro/mdro_5.html. Updated December 29, 2009. Accessed November 2014.
- 6. Bearman GM, Marra AR, Sessler CN, et al. A controlled trial of universal gloving versus contact precautions for preventing the transmission of multidrug-resistant organisms. *Am J Infect Control* 2007;35:650–655.
- Safdar N, Marx J, Meyer NA, Maki DG. Effectiveness of preemptive barrier precautions in controlling nosocomial colonization and infection by methicillin-resistant *Staphylococcus aureus* in a burn unit. *Am J Infect Control* 2006;34:476–483.
- Trick WE, Weinstein RA, DeMarais PL, et al. Comparison of routine glove use and contact-isolation precautions to prevent transmission of multidrug-resistant bacteria in a long-term care facility. J Am Geriatr Soc 2004;52:2003–2009.
- 9. Harbarth S, Fankhauser C, Schrenzel J, et al. Universal screening for methicillin-resistant *Staphylococcus aureus* at hospital admission and nosocomial infection in surgical patients. *JAMA* 2008;299:1149–1157.
- 10. Marshall C, Richards M, McBryde E. Do active surveillance and contact precautions reduce MRSA acquisition? A prospective interrupted time series. *PLOS ONE* 2013;8:e58112.
- 11. Derde LP, Cooper BS, Goossens H, et al. Interventions to reduce colonisation and transmission of antimicrobial-resistant bacteria in intensive care units: an interrupted time series study and cluster randomised trial. *Lancet Infect Dis* 2014;14:31–39.
- Huskins WC, Huckabee CM, O'Grady NP, et al. Intervention to reduce transmission of resistant bacteria in intensive care. *N Engl J Med* 2011;364:1407–1418.
- Huang SS, Yokoe DS, Hinrichsen VL, et al. Impact of routine intensive care unit surveillance cultures and resultant barrier precautions on hospital-wide methicillin-resistant *Staphylococcus aureus* bacteremia. *Clin Infect Dis* 2006;43:971–978.
- Lucet JC, Paoletti X, Lolom I, et al. Successful long-term program for controlling methicillin-resistant *Staphylococcus aureus* in intensive care units. *Intensive Care Med* 2005;31: 1051–1057.
- Robicsek A, Beaumont JL, Paule SM, et al. Universal surveillance for methicillin-resistant *Staphylococcus aureus* in 3 affiliated hospitals. *Ann Intern Med* 2008;148:409–418.
- Harris AD, Pineles L, Belton B, et al. Universal glove and gown use and acquisition of antibiotic-resistant bacteria in the ICU: a randomized trial. *JAMA* 2013;310:1571–1580.
- Jain R, Kralovic SM, Evans ME, et al. Veterans Affairs initiative to prevent methicillin-resistant *Staphylococcus aureus* infections. *N Engl J Med* 2011;364:1419–1430.
- Fournier S, Monteil C, Lepainteur M, et al. Long-term control of carbapenemase-producing Enterobacteriaceae at the scale of a large French multihospital institution, a nine-year experience, France, 2004 to 2012. *Euro Surveill* 2014;19. pii:20802.
- Rossini FA, Fagnani R, Leichsenring ML, et al. Successful prevention of the transmission of vancomycin-resistant enterococci in a Brazilian public teaching hospital. *Rev Soc Bras Med Trop* 2012;45:184–188.

- Carmona F, Prado SI, Silva MF, et al. Vancomycin-resistant *Enterococcus* outbreak in a pediatric intensive care unit: report of successful interventions for control and prevention. *Braz J Med Biol Res* 2012;45:158–162.
- Rosenberger LH, Hranjec T, Politano AD, et al. Effective cohorting and "superisolation" in a single intensive care unit in response to an outbreak of diverse multi-drug-resistant organisms. Surg Infect (Larchmt) 2011;12:345–350.
- 22. Xu HT, Tian R, Chen DK, et al. Nosocomial spread of hospitaladapted CC17 vancomycin-resistant *Enterococcus faecium* in a tertiary-care hospital of Beijing, China. *Chin Med J (Engl)* 2011;124:498–503.
- 23. Ergaz Z, Arad I, Bar-Oz B, et al. Elimination of vancomycinresistant enterococci from a neonatal intensive care unit following an outbreak. *J Hosp Infect* 2010;74:370–376.
- Servais A, Mercadal L, Brossier F, et al. Rapid curbing of a vancomycin-resistant *Enterococcus faecium* outbreak in a nephrology department. *Clin J Am Soc Nephrol* 2009;4: 1559–1564.
- Nolan SM, Gerber JS, Zaoutis T, et al. Outbreak of vancomycinresistant *Enterococcus* colonization among pediatric oncology patients. *Infect Control Hosp Epidemiol* 2009;30:338–345.
- Cheng VC, Chan JF, Tai JW, et al. Successful control of vancomycin-resistant *Enterococcus faecium* outbreak in a neurosurgical unit at non-endemic region. *Emerg Health Threats J* 2009;2:e9.
- 27. Hoshuyama T, Moriguchi H, Muratani T, Matsumoto T. Vancomycin-resistant enterococci (VRE) outbreak at a university hospital in Kitakyushu, Japan: case-control studies. *J Infect Chemother* 2008;14:354–360.
- Aumeran C, Baud O, Lesens O, Delmas J, Souweine B, Traoré O. Successful control of a hospital-wide vancomycin-resistant *Enterococcus faecium* outbreak in France. *Eur J Clin Microbiol Infect Dis* 2008;27:1061–1064.
- Schmidt-Hieber M, Blau IW, Schwartz S, et al. Intensified strategies to control vancomycin-resistant enterococci in immunocompromised patients. *Int J Hematol* 2007;86:158–162.
- Lucet JC, Armand-Lefevre L, Laurichesse JJ, et al. Rapid control of an outbreak of vancomycin-resistant enterococci in a French university hospital. *J Hosp Infect* 2007;67:42–48.
- Yoonchang SW, Peck KR, Kim OS, et al. Efficacy of infection control strategies to reduce transmission of vancomycin-resistant enterococci in a tertiary care hospital in Korea: a 4-year followup study. *Infect Control Hosp Epidemiol* 2007;28:493–495.
- 32. Wang JT, Chen YC, Chang SC, et al. Control of vancomycinresistant enterococci in a hospital: a five-year experience in a Taiwanese teaching hospital. *J Hosp Infect* 2004;58:97–103.
- Hachem R, Graviss L, Hanna H, et al. Impact of surveillance for vancomycin-resistant enterococci on controlling a bloodstream outbreak among patients with hematologic malignancy. *Infect Control Hosp Epidemiol* 2004;25:391–394.
- 34. Bearman G, Rosato AE, Duane TM, et al. Trial of universal gloving with emollient-impregnated gloves to promote skin health and prevent the transmission of multidrug-resistant organisms in a surgical intensive care unit. *Infect Control Hosp Epidemiol* 2010;31:491–497.
- 35. De Angelis G, Cataldo MA, De Waure C, et al. Infection control and prevention measures to reduce the spread of vancomycinresistant enterococci in hospitalized patients: a systematic review and meta-analysis. J Antimicrob Chemother 2014;69:1185–1192.

- 36. 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.
- 37. Evans HL, Shaffer MM, Hughes MG, et al. Contact isolation in surgical patients: a barrier to care? *Surgery* 2003;134:180–188.
- Kirkland KB, Weinstein JM. Adverse effects of contact isolation. *Lancet* 1999;354:1177–1178.
- 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.
- Masse V, Valiquette L, Boukhoudmi S, et al. Impact of methicillin resistant *Staphylococcus aureus* contact isolation units on medical care. *PLOS ONE* 2013;8:e57057.
- Saint S, Higgins LA, Nallamothu BK, Chenoweth C. Do physicians examine patients in contact isolation less frequently? A brief report. *Am J Infect Control* 2003;31:354–356.
- 42. Gilligan P, Quirke M, Winder S, Humphreys H. Impact of admission screening for methicillin-resistant *Staphylococcus aureus* on the length of stay in an emergency department. *J Hosp Infect* 2010;75:99–102.
- 43. McLemore A, Bearman G, Edmond MB. Effect of contact precautions on wait time from emergency room disposition to inpatient admission. *Infect Control Hosp Epidemiol* 2011;32:298–299.
- 44. Goldszer RC, Tamplin E, Yokoe DS, et al. A program to remove patients from unnecessary contact precautions. *J Clin Outcomes Manag* 2002;9:553–556.
- 45. Stelfox HT, Bates DW, Redelmeier DA. Safety of patients isolated for infection control. *JAMA* 2003;290:1899–1905.
- 46. Karki S, Leder K, Cheng AC. Patients under contact precautions have an increased risk of injuries and medication errors: a retrospective cohort study. *Infect Control Hosp Epidemiol* 2013;34:1118–1120.
- 47. Catalano G, Houston SH, Catalano MC, et al. Anxiety and depression in hospitalized patients in resistant organism isolation. *South Med J* 2003;96:141–145.
- Davies H, Rees J. Psychological effects of isolation nursing, 1: mood disturbance. *Nurs Stand* 2000;14:35–38.
- Rees J, Davies HR, Birchall C, Price J. Psychological effects of source isolation nursing, 2: patient satisfaction. *Nurs Stand* 2000;14:32–36.
- Day HR, Perencevich EN, Harris AD, et al. Association between contact precautions and delirium at a tertiary care center. *Infect Control Hosp Epidemiol* 2012;33:34–39.
- Gammon J. Analysis of the stressful effects of hospitalisation and source isolation on coping and psychological constructs. *Int J Nurs Pract* 1998;4:84–96.
- Kennedy P, Hamilton LR. Psychological impact of the management of methicillin-resistant *Staphylococcus aureus* (MRSA) in patients with spinal cord injury. *Spinal Cord* 1997;35:617–619.
- Day HR, Morgan DJ, Himelhoch S, Young A, Perencevich EN. Association between depression and contact precautions in veterans at hospital admission. *Am J Infect Control* 2011;39:163–165.
- 54. Knowles HE. The experience of infectious patients in isolation. *Nurs Times* 1993;89:53–56.
- Tarzi S, Kennedy P, Stone S, Evans M. Methicillin-resistant Staphylococcus aureus: psychological impact of hospitalization and isolation in an older adult population. J Hosp Infect 2001;49:250–254.
- 56. Day HR, Perencevich EN, Harris AD, et al. Do contact precautions cause depression? A two-year study at a tertiary care medical centre. *J Hosp Infect* 2011;79:103–107.

- 57. Cohen E, Austin J, Weinstein M, Matlow A, Redelmeier DA. Care of children isolated for infection control: a prospective observational cohort study. *Pediatrics* 2008;122:e411–e415.
- Mehrotra P, Croft L, Day HR, et al. Effects of contact precautions on patient perception of care and satisfaction: a prospective cohort study. *Infect Control Hosp Epidemiol* 2013;34:1087–1093.
- 59. Gasink LB, Singer K, Fishman NO, et al. Contact isolation for infection control in hospitalized patients: is patient satisfaction affected? *Infect Control Hosp Epidemiol* 2008;29:275–278.
- Siegel JD, Rhinehart E, Jackson M, Chiarello L, Healthcare Infection Control Practices Advisory Committee. Management of multidrug-resistant organisms in health care settings, 2006. *Am J Infect Control* 2007;35:S165–S193.
- 61. Dhar S, Marchaim D, Tansek R, et al. Contact precautions: more is not necessarily better. *Infect Control Hosp Epidemiol* 2014;35:213–221.
- 62. Shenoy ES, Kim J, Rosenberg ES, et al. Discontinuation of contact precautions for methicillin-resistant *Staphylococcus aureus*: a randomized controlled trial comparing passive and active screening with culture and polymerase chain reaction. *Clin Infect Dis* 2013;57:176–184.

- Edmond M. Panel on clinical controversies in ID. In: Program and abstracts of IDWeek 2014; Philadelphia, PA; October 9, 2014. Abstract 18. https://idsa.confex.com/idsa/2014/webprogram/ Session6312.html. Accessed November 2014.
- 64. Gandra S, Barysauskas CM, Mack DA, Barton B, Finberg R, Ellison RT. Impact of contact precautions on falls, pressure ulcers and transmission of MRSA and VRE in hospitalized patients. *J Hosp Infect* 2014;88:170–176.
- 65. Haessler S. MRSA success stories. In *Proceedings of SHEA Spring Conference 2014*. Denver, CO: SHEA, 2014.
- 66. Kirkland KB. Taking off the gloves: toward a less dogmatic approach to the use of contact isolation. *Clin Infect Dis* 2009;48:766–771.
- 67. Fraser TG, Fatica C, Scarpelli M, et al. Decrease in *Staphylococcus aureus* colonization and hospital-acquired infection in a medical intensive care unit after institution of an active surveillance and decolonization program. *Infect Control Hosp Epidemiol* 2010;31:779–783.
- Derde LP, Dautzenberg MJ, Bonten MJ. Chlorhexidine body washing to control antimicrobial-resistant bacteria in intensive care units: a systematic review. *Intensive Care Med* 2012;38:931–939.