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Original Article

Assessment of antibiotic-resistant organism transmission among rooms of hospitalized patients, healthcare personnel, and the hospital environment utilizing surrogate markers and selective bacterial cultures

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Abstract

Objective: To assess potential transmission of antibiotic-resistant organisms (AROs) using surrogate markers and bacterial cultures.

Design: Pilot study.

Setting: A 1,260-bed tertiary-care academic medical center.

Participants: The study included 25 patients (17 of whom were on contact precautions for AROs) and 77 healthcare personnel (HCP).

Methods: Fluorescent powder (FP) and MS2 bacteriophage were applied in patient rooms. HCP visits to each room were observed for 2–4 hours; hand hygiene (HH) compliance was recorded. Surfaces inside and outside the room and HCP skin and clothing were assessed for fluorescence, and swabs were collected for MS2 detection by polymerase chain reaction (PCR) and selective bacterial cultures.

Results: Transfer of FP was observed for 20 rooms (80%) and 26 HCP (34%). Transfer of MS2 was detected for 10 rooms (40%) and 15 HCP (19%). Bacterial cultures were positive for 1 room and 8 HCP (10%). Interactions with patients on contact precautions resulted in fewer FP detections than interactions with patients not on precautions ($P < .001$); MS2 detections did not differ by patient isolation status. Fluorescent powder detections did not differ by HCP type, but MS2 was recovered more frequently from physicians than from nurses ($P = .03$). Overall, HH compliance was better among HCP caring for patients on contact precautions than among HCP caring for patients not on precautions ($P = .003$), among nurses than among other nonphysician HCP at room entry ($P = .002$), and among nurses than among physicians at room exit ($P = .03$). Moreover, HCP who performed HH prior to assessment had fewer fluorescence detections ($P = .008$).

Conclusions: Contact precautions were associated with greater HCP HH compliance and reduced detection of FP and MS2.

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Antibiotic-resistant organisms (AROs) present a major infection control threat for patients in hospitals, and they increase the risk of serious healthcare-associated infections. Hospital environmental surfaces can become contaminated with AROs and may contribute to ARO transmission, either directly or via the hands or clothing of healthcare personnel (HCP).^{1–5} Contact precautions (gowns and gloves) have been an essential component of infection

prevention practices to limit transmission of AROs.⁶ However, there has been debate regarding whether contact precautions are effective in reducing ARO transmission.^{7–9}

The relationships among environmental contamination, HCP cross contamination, and ARO transmission are difficult to study. Previous studies have demonstrated that contaminated hospital surfaces can contribute to the spread of nosocomial infections.^{10–12} Other studies have demonstrated ARO transfer from infected patients or contaminated surfaces to the hands and clothing of HCP.^{13–17} However, few studies have focused on the relationship between contaminated surfaces in patient rooms and the risk of HCP cross contamination outside patient rooms.¹⁸

Additional studies that examine the associations between environmental surface contamination, HCP cross contamination and

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ARO transmission patterns, and the impact of contact isolation practices on these associations, are needed to better inform policies and procedures for the use of personal protective equipment (PPE) and to reduce ARO transmission and healthcare-associated infections (HAIs). Surrogate markers, such as fluorescent powder (FP) and MS2, a nonpathogenic bacteriophage, are unique tools for studying ARO transmission and cross contamination in hospitals.^{19,20} Fluorescent powder and MS2 have been used to study HCP self-contamination while donning/doffing PPE^{17,21,22} and the effectiveness of hospital cleaning procedures.^{23–25}

The aim of this prospective cohort study was to assess ARO transmission and cross contamination patterns in real-world hospital settings using 2 surrogate markers (FP and MS2 bacteriophage) and selective bacterial cultures.

Methods

This study was conducted in a general medicine ward, a medical intensive care unit (ICU), and an emergency department (ED) at a 1,260-bed tertiary-care academic hospital in St Louis, Missouri. Patients aged ≥ 18 years hospitalized between September 16, 2015, and February 9, 2016, as well as HCP caring for the enrolled patients were eligible for inclusion. The study protocol was reviewed and approved by the Washington University Human Research Protection Office. Written informed consent was obtained from all patients or a legally authorized representative. Participating HCP provided verbal consent prior to study participation.

Patient enrollment

Two patients on contact precautions for vancomycin-resistant *Enterococcus* (VRE) or methicillin-resistant *Staphylococcus aureus* (MRSA) were enrolled for each patient not on contact precautions. At enrollment, each patient's room was scanned for fluorescence using an ultraviolet (UV) light. If fluorescence was detected, the area was wiped clean before surrogate marker application. For patients on contact precautions, flocked swab collection kits (ESwab, Copan Diagnostics, Murietta, CA) were used to collect swabs from each of the surfaces targeted for surrogate marker application and to collect nasal, axilla, inguinal, and stool or rectal swabs from each patient. Baseline patient and environmental samples were determined using selective bacterial culture.

Surrogate marker application

In each patient room, 4 high-touch surfaces were selected for surrogate marker application: the front of the patient's gown, the top of each bed rail, and the bedside table or computer mouse. Fluorescent powder (0.02 g, Glo Germ, Moab, UT) was applied to each surface using a brush applicator. MS2 bacteriophage (1:10 dilution of commercially available stock solution in viral transport medium, 1.0×10^8 PFU/mL per site,¹⁷ ZeptoMetrix, Buffalo, NY) was applied using an atomizer (Teleflex, Morrisville, NC).

HCP enrollment and observations

Following surrogate marker application, trained study coordinators observed each patient room for 2–4 hours from the hallway. During this period, HCP hand hygiene (HH) compliance at room entry and exit, defined as the use of alcohol hand rub or soap and water, were recorded, and the first 3 surfaces that each HCP touched after exiting the room were flagged for later assessment. Also, 3–4 HCP who entered the room during the observation period were recruited for study participation.

Sample collection

After the first visit to a patient's room, participating HCP had their hands, face and hair, and clothing scanned with a UV light to identify areas of fluorescence. For patients on contact precautions, UV scanning was performed after the HCP had removed PPE. HCP were assessed only once, even if they visited the room multiple times. At the end of the observation period, the patient's room, the first 3 surfaces that each participating HCP had touched after exiting the room, and 4 additional locations on the study ward (ie, medication cabinet, door handles, nurse's station, and elevator buttons) were scanned for fluorescence.

Areas that fluoresced were photographed and trained study coordinators collected surface samples using a viral transport collection kit (Quidel, San Diego, CA). Additional samples were collected from the 4 locations on the study ward and from each participating HCP hands and gloves, face (ie, periorbital, nasal, and oral areas), and sleeve and wrist. These samples were tested for the presence of MS2.

If the patient was on contact precautions, flocked swab collection kits were used to collect additional samples from each area where fluorescence was observed and from the 4 selected locations on the study ward. One pooled sample was also collected from the face, hands, and wrists of each participating HCP. These swabs were submitted for selective bacterial culture.

After sample collection, the surfaces where the surrogate markers had been applied and any areas where fluorescence was observed were wiped clean to prevent further transmission of FP and MS2. Each patient room was used only once to further minimize the possibility of residual marker from a previous patient.

Bacterial culture

Swabs collected to identify MS2 contamination had RNA extracted from the transport medium using the QIAamp viral RNA Mini Kit (Qiagen, Germantown, MD). Real-time reverse transcriptase polymerase chain reaction (PCR) was used to detect MS2 bacteriophage using the Cepheid Smart Cyclex (Cepheid, Sunnyvale, CA).

Swabs associated with patients on contact precautions were cultured for VRE, MRSA, and methicillin-susceptible *Staphylococcus aureus* (MSSA). Swabs were plated to CHROMID VRE chromogenic medium (bioMérieux, Marcy-l'Étoile, France) to select for VRE; on Spectra MRSA chromogenic agar (Remel, Lenexa, KS) to select for MRSA; and on 5% sheep's blood agar (Hardy Diagnostics, Santa Maria, CA) to recover MSSA. All swabs were also inoculated to 6.5% NaCl broth (Hardy Diagnostics) as an enrichment method to recover VRE, MRSA, and MSSA if these did not grow on the primary plated media. When growth was observed, 4 colonies of each type of organism were subcultured and identified using matrix-assisted laser desorption ionization time-of-flight mass spectrometry (MALDI-TOF MS) using VITEK MS.^{26–28} After bacterial identification was confirmed, phenotypic antimicrobial susceptibility testing and repetitive sequence-based PCR (repPCR) were performed. Staphylococcal cassette chromosome *mec* (SCC*mec*) typing was performed on all *S. aureus* isolates.^{29,30}

Statistical analysis

Patterns in the location and type of surrogate marker detections were evaluated qualitatively. Odds ratios with 95% confidence intervals and the χ^2 or Fisher exact test were used to characterize associations between predictor and outcome variables. Predictor

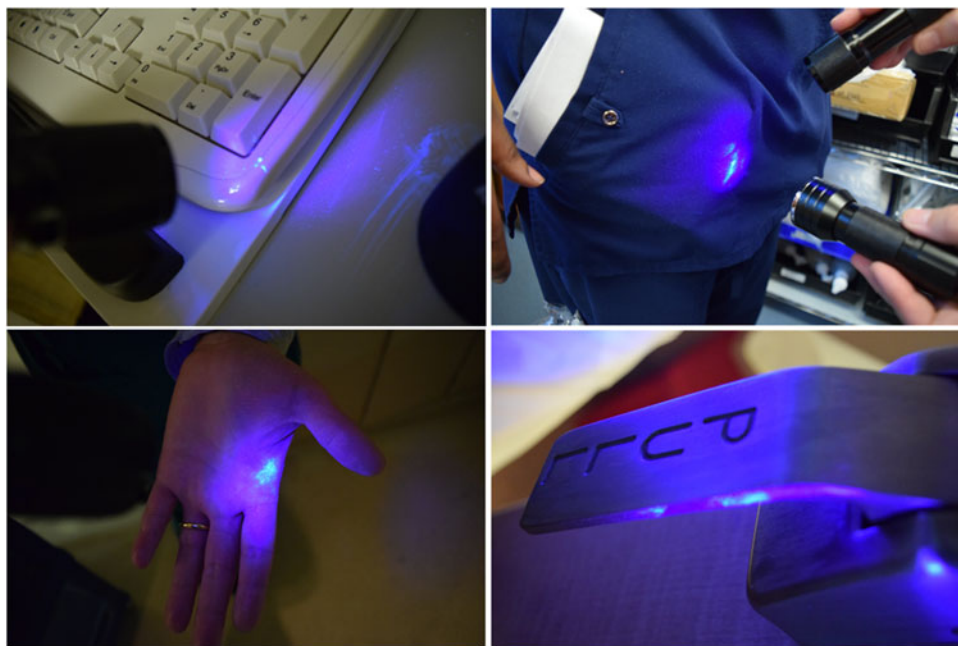


Fig. 1. Examples of fluorescent powder (FP) detections observed in this study.

variables included patient contact isolation status and type of HCP. Outcome variables included FP, MS2, and VRE, MRSA, or MSSA detections in patient rooms, on HCP, and/or on surfaces touched by HCP. The use of HH by HCP at room entry and exit were assessed as both predictor and outcome variables. Two measures of HCP HH compliance were examined: (1) HH at the first room visit by participating HCP and (2) HH over all room visits by all HCP. The first measure was used to determine the association between HH and surrogate marker detections, and the second provided a more complete picture of HH practices overall. All analyses were performed using SPSS version 24.0 software (IBM, Armonk, NY).

Results

In total, 25 patients were enrolled: 10 in the medicine ward, 10 in the ICU, and 5 in the ED. Among them, 17 patients (68%) were on contact precautions for VRE ($n = 12$), MRSA ($n = 4$), or VRE and MRSA ($n = 1$). In addition, 77 HCP participated in the study: half ($n = 40$, 52%) were nurses ($n = 35$), nurse practitioners ($n = 3$), or student nurses ($n = 2$). Other participating HCP included physicians ($n = 16$, 21%), patient care technicians ($n = 9$, 12%), respiratory therapists ($n = 4$, 5%), radiology technicians ($n = 2$, 3%), dietitians ($n = 2$, 3%), 1 pharmacist (1%), a pharmacy student, an infection preventionist, and a unit secretary.

Fluorescent powder detections

In 20 patient rooms (80%), fluorescence was detected on at least 1 site outside the areas where FP had been applied, most commonly on the computer keyboard ($n = 15$), the counter ($n = 7$), or the door handle ($n = 5$). In 3 cases, fluorescence was also detected in the study ward, at the nurses' station ($n = 2$) or on the medication cabinet ($n = 1$). Moreover, fluorescence was detected on 26 HCP (34%): on their body, hands, or clothing ($n = 23$) and/or on a surface they touched after exiting the patient's room ($n = 10$). Examples of FP detections are shown in Figure 1.

The HCP caring for patients on contact precautions had significantly fewer FP detections, on themselves and/or on the surfaces

they touched, than HCP caring for patients not on precautions (19% vs 70%; $P < .001$) (Table 1). We found no significant difference in the rates of FP detection among different types of HCP (Table 2).

MS2 detections

MS2 was detected in 9 patient rooms (36%), most commonly on the computer ($n = 4$) and, outside 1 room, on a medication cabinet. Moreover, 15 HCP (19%) had MS2 detections, either on their body or clothing ($n = 10$) and/or on surfaces touched after exiting a patient room ($n = 6$), most commonly the door handle ($n = 3$). Also, 1 HCP had MS2 identified on 2 sites on the body or clothing, and 1 HCP had MS2 identified on 2 touched surfaces.

In general, MS2 was recovered less frequently on HCP and/or surfaces touched by HCP caring for patients on contact precautions than on HCP caring for patients not on precautions, but these differences did not reach statistical significance (Table 1). MS2 was more often detected on physicians than on nurses (40% vs 27%; $P = .02$) (Table 2).

Bacterial culture results

Of the patients on contact precautions, 12 (71%) had baseline swabs positive for the ARO for which the patient was placed on contact precautions. Also, 2 patients, 1 on precautions for MRSA and 1 for VRE, had swabs positive for both MRSA and VRE. One patient on precautions for VRE had baseline swabs positive for MSSA.

Moreover, 7 patients on contact precautions (41%) had 1 or more room surfaces with a positive baseline bacterial culture (Table 3). For 6 patients, the organism identified was the organism that triggered contact precautions; 2 patients had surfaces that were also positive for MSSA. The remaining patient, who was on precautions for VRE but had a baseline swab positive for MSSA, had baseline room surface swabs that were also positive for MSSA.

Among the swabs collected from surfaces where fluorescence was observed, only 2 had a positive bacterial culture (Table 3).

Table 1. Fluorescent Powder and MS2 Detections on Participating Healthcare Personnel (HCP) and Surfaces Touched by Participating HCP After Exiting the Patient's Room, by Patient Isolation Status

Type of Detection	All HCP (N = 77), No. (%)	HCP Caring for Patient on Contact Precautions (N = 54), No. (%)	HCP Caring for Patient Not on Precautions (N = 23), No. (%)	OR (95% CI)	P Value ^a
Fluorescent powder^b	26 (34)	10 (19)	16 (70)	0.10 (0.03–0.31)	<.001
HCP ^c	23 (30)	9 (17)	14 (61)	0.13 (0.04–0.39)	<.001
Touched surface ^d	10 (13)	3 (6)	7 (30)	0.13 (0.03–0.58)	.006
MS2^e	15 (19)	8 (15)	7 (30)	0.40 (0.12–1.27)	.13
HCP ^c	10 (13)	4 (7)	6 (26)	0.23 (0.06–0.90)	.06
Touched surface ^d	6 (8)	4 (7)	2 (9)	0.84 (0.14–4.94)	1.00

Note. OR, odds ratio; CI, confidence interval; PCR, polymerase chain reaction.

^aFisher's exact test was used for comparisons due to small cell sizes.

^bDefined as the visualization of fluorescence when the HCP or surface was scanned with a handheld UV light.

^cIncludes HCP hands, sleeves/wrist, gloves, face, and clothing.

^dEnvironmental surfaces touched by HCP after leaving the patient room.

^eDefined as the detection of MS2 on a swab collected from the HCP or surface via real-time reverse-transcriptase PCR.

Table 2. Fluorescent Powder and MS2 Detections on Participating Healthcare Personnel (HCP) and/or Surfaces Touched by HCP Type

Type of Detection	Surrogate Marker Detected, No. (%)	Surrogate Marker Not Detected, No. (%)	OR (95% CI)	P Value
Fluorescent powder	N = 26	N = 51		
Nurse (n = 40) ^a	13 (50)	26 (51)	Reference	
Physician (n = 16)	3 (12)	13 (26)	0.48 (0.12–1.98)	.31
Other (n = 21) ^b	10 (39)	12 (24)	1.89 (0.64–5.57)	.25
MS2	N = 15	N = 62		
Nurse (n = 40) ^a	4 (27)	35 (57)	Reference	
Physician (n = 16)	6 (40)	10 (16)	5.40 (1.27–22.93)	.02
Other (n = 21) ^b	5 (33)	17 (27)	2.57 (0.67–11.88)	.16

Note. OR, odds ratio; CI, confidence interval.

^aIncludes nurse practitioners and student nurses.

^bIncludes patient care technicians, respiratory therapists, radiology techs, dieticians, pharmacist, pharmacy student, infection prevention technician, and unit secretary.

One sample, from the foot of a bed, was positive for VRE. The other, from an elevator button, was positive for MSSA. Both were associated with the same patient, who had a baseline swab positive for VRE.

Of the 54 HCP who cared for a patient on contact precautions, 8 (15%) had a positive pooled swab, all of which were positive for MSSA (Table 3). These HCP had cared for 4 different patients, none of whom had a baseline swab positive for MSSA, but one of whom had a baseline room surface positive for MSSA.

Only 2 HCP (4%) who cared for a patient on contact precautions had a touched surface with a positive bacterial culture (Table 3). The first, a blood glucose monitor, was positive for VRE, although the HCP was positive for MSSA, and MRSA was identified in the patient's room. The second, a door handle, was positive for MSSA and was touched by an HCP who was also positive for MSSA, although VRE was identified in the patient's room.

Among samples that were positive for VRE, 6 strain types were identified by repPCR. The most common, type C, was associated with 8 patients, 3 of whom were also positive for type B. Among these, 2 patients had VRE type C identified in both patient and room-surface samples. Another patient had multiple VRE types (A, D, E, F) identified in patient and room-surface samples.

Among samples that were positive for *S. aureus*, 9 strain types were identified by repPCR (3 among MRSA samples and 8 among MSSA samples); 4 strain types were identified by SCCmec typing. Also, 4 of the 5 patients who were positive for MRSA and 2 of the 3 patients with room surfaces positive for MRSA had the same strain typing (repPCR B, SCCmec IV). Among 8 HCP who were positive for MSSA, 7 had the same SCCmec type (III). Of these samples, 3 were repPCR type F; the others had diverse repPCR typing.

HCP hand hygiene observations

Both measures of HCP HH compliance yielded similar estimates. HH compliance was lower at room entry than at room exit. Only 18% of HCP performed HH at room entry (14 of 77 first visits by participating HCP and 54 of 298 total HCP visits), and 52% performed HH at room exit (40 of 77 first visits and 54 of 290 total visits).

The HCP HH compliance at room entry did not differ by patient isolation status (Table 4). However, compliance at room exit was better among the HCP caring for patients on contact precautions than among HCP caring for patients not on precautions: 61% versus 30% first room visits ($P = .02$) and 58% versus 37% all room visits ($P < .01$). We observed no differences in HH

Table 3. Microbiologic Culture Results for the Patients on Contact Precautions and the Healthcare Personnel (HCP) Who Cared for These Patients

Samples	All Positive	VRE	MRSA	MSSA ^a
Samples from patients (n = 17)				
Baseline patient swabs ^b	13 ^c	9 ^d	5 ^e	1 ^f
Baseline room surface swabs	7 ^g	3 ^h	3 ⁱ	3 ^j
Surface swabs from areas where fluorescence was observed inside patient rooms	1	1 ^k	0	0
Surface swabs from areas where fluorescence was observed outside patient rooms	1	0	0	1 ^l
Samples from HCP (n = 54)				
Pooled swab from face, hands, and wrist	8	0	0	8 ^m
Swabs collected from surfaces touched by participating HCP after leaving patient rooms	2	1 ⁿ	0	1 ^o

Note. VRE, vancomycin-resistant *Enterococcus*; MRSA, methicillin-resistant *Staphylococcus aureus*; MSSA, methicillin-susceptible *Staphylococcus aureus*; NA, not applicable.

^aMSSA is not an indication for contact precautions.

^bIncluded nasal, axilla, inguinal skin, and stool or rectal swabs.

^cFor 12 patients, the identified organism matched the reason for contact precautions; 2 patients had swabs that were positive for both VRE and MRSA; 1 patient had swabs that were positive for only MSSA.

^dRepetitive sequence-based PCR (repPCR) results were variable: samples from 1 patient were type B; samples from 4 patients were type C; samples from 2 patients were types B and C; samples from 1 patient were types A, B, and C; and samples from 1 patient were types A, D, E, and F.

^eSamples from 4 patients were repetitive sequence-based PCR (repPCR) type B and staphylococcal cassette chromosome *mec* (SCC*mec*) type IV. Samples from 1 patient were repPCR types E and G, SCC*mec* type III.

^fSamples were repPCR types C and B, SCC*mec* type III.

^gFor 6 patients, the identified organism matched the reason for contact precautions; 1 patient had baseline room surface swabs that were positive for both MRSA and MSSA; and 1 patient had swabs that were positive for both VRE and MSSA. One patient had baseline room-surface swabs that were positive for only MSSA.

^hBaseline room-surface samples from 2 patient rooms were all repPCR type C; samples from the third room were repPCR types D and F.

ⁱSamples from 2 rooms were repPCR type B, SCC*mec* type IV. Samples from one room were repPCR types E and G, SCC*mec* type III.

^jSamples from the first room were repPCR type B, SCC*mec* type III. Samples from the second room were repPCR type E, SCC*mec* type I. Samples from the third room were repPCR types B and D, SCC*mec* type III.

^kSamples were repPCR types D and E.

^lThese samples were repPCR type A, SCC*mec* type I.

^mOne HCP had samples that were repPCR type A, SCC*mec* type III; 1 HCP had samples that were repPCR type B, SCC*mec* type III; 1 HCP had samples that were repPCR type E, SCC*mec* type I; 3 HCP had samples that were repPCR type F, SCC*mec* type III; 1 HCP had samples that were repPCR types A, C, and D, SCC*mec* type III; and 1 HCP had samples that were repPCR types E and D, SCC*mec* type III.

ⁿSamples were repPCR type C.

^oTwo surfaces touched by the same HCP had samples that were positive for MSSA types H and A, SCC*mec* type III.

Table 4. Healthcare Personnel (HCP) Observations Where Hand Hygiene Was Performed, at First Room Entry/Exit and All Room Entries/Exits, by Patient Isolation Status

Observation	Patients on Contact Precautions, No. (%)	Patients Not on Contact Precautions, No. (%)	OR (95% CI)	P Value ^a
First room visit by participating HCP				
Room entry	N = 54 8 (15)	N = 23 6 (26)	2.03 (0.61–6.71)	.33
Room exit	33 (61)	7 (30)	0.28 (0.10–0.79)	.02
All HCP room visits				
Room entry	N = 221 38 (17)	N = 77 16 (21)	1.26 (0.66–2.43)	.50
Room exit ^b	124 (58)	28 (37)	0.44 (0.26–0.75)	.003

Note. OR, odds ratio; CI, confidence interval.

^aThe Fisher exact test was used for comparisons due to small cell sizes.

^b8 room-exit observations were missing because room exit could not be observed.

compliance at first room visit for nurses versus physicians or other HCP (Table 5). However, when considering all room visits, nurses were more likely than other nonphysician HCP to perform HH at room entry (25% vs 8%; $P < .01$) and were more likely than physicians to perform HH at room exit (59% vs 43%; $P = .03$).

The association between HCP HH and surrogate marker detections is shown in Table 6. Although few associations were observed between either HH measure and surrogate marker detections, HCP who performed HH immediately after the first room exit and before being swabbed were less likely than HCP who did not perform HH at room exit to have fluorescence detected (20% vs 49%; $P = .008$).

Discussion

In this study, the transfer of both FP and MS2 were observed both inside and outside patient rooms, on participating HCP, and on surfaces touched by HCP after exiting patient rooms. Transfer of FP occurred more frequently than the transfer of MS2, and positive bacterial cultures were even less frequent.

Although few studies have utilized both FP and MS2 as surrogate markers, some have also reported higher rates of FP compared to MS2 detections.^{17,31} Others have reported similar detection rates^{19,32} or more frequent MS2 detections.²¹ This lack of agreement may indicate that neither marker performs significantly

Table 5. Healthcare Personnel (HCP) Hand Hygiene Observations, at First Room Entry/Exit and All Room Entries/Exits, by HCP Type

Observation	Hand Hygiene Performed, No (%)	OR (95% CI)	P Value
First room visit by participating HCP			
Room entry (n = 77)			
Nurse (n = 40) ^a	7 (18)	Reference	
Physician (n = 16)	5 (31)	0.47 (0.12–1.77)	.26
Other (n = 21) ^b	2 (10)	2.02 (0.38–10.70)	.41
Room exit (n = 77)			
Nurse (n = 40) ^a	19 (48)	Reference	
Physician (n = 16)	9 (56)	0.70 (0.22–2.26)	.56
Other (n = 21) ^b	12 (57)	0.68 (0.23–1.97)	.48
All HCP room visits			
Room entry (n = 298)			
Nurse (n = 150) ^a	38 (25)	Reference	
Physician (n = 71)	10 (14)	2.07 (0.97–4.44)	.06
Other (n = 77) ^b	6 (8)	4.02 (1.62–9.98)	.003
Room exit^c (n = 290)			
Nurse (n = 147) ^a	87 (59)	Reference	
Physician (n = 68)	29 (43)	1.95 (1.09–3.49)	.03
Other (n = 75) ^b	36 (48)	1.57 (0.90–2.75)	.11

Note. OR, odds ratio; CI, confidence interval.

^aIncludes student nurses and nurse practitioners.

^bIncludes patient care technicians, respiratory therapists, radiology techs, dieticians, pharmacist, pharmacy student, infection prevention technician, and unit secretary.

^c8 room-exit observations were missing because room exit could not be observed.

Table 6. Association Between Hand Hygiene Performance at Room Exit and Detection of Fluorescence and MS2 on Healthcare Personnel (HCP) and on Environmental Surfaces Touched by HCP

Observation	Hand Hygiene Performed, No. (%)	Hand Hygiene Not Performed, No. (%)	OR (95% CI)	P Value
First room exit by participating HCP		N = 41	N = 36	
Fluorescent powder detected	8 (20)	18 (49)	3.79 (1.38–10.38)	.008
MS2 detected	8 (20)	7 (19)	0.93 (0.30–2.89)	.91
All HCP room exits		N = 29	N = 48	
Fluorescent powder detected	7 (24)	19 (40)	2.06 (0.74–5.76)	.17
MS2 detected	7 (24)	8 (17)	0.63 (0.20–1.97)	.42

Note. OR, odds ratio; CI, confidence interval.

better than the other, or it may be related to differences in the means of detection (visual versus swabs). However, the 2 markers are thought to model different types of contamination: FP may model gross bacterial contamination and MS2 may simulate viral contamination events.²¹ Therefore, different detection rates may be reasonable. More data are needed to determine which surrogate markers are better models for the study of ARO transmission.

In contrast to surrogate markers, bacterial culture may identify actual ARO transmission events. This study focused on 2 AROs that routinely trigger contact precautions, MRSA and VRE, as well as MSSA. Although MSSA does not routinely trigger contact precautions, it is a clinically relevant pathogen that causes significant morbidity in hospitalized patients.^{33,34} The greater frequency of surrogate marker detections as compared to ARO detections may suggest that FP and MS2 overrepresent the likelihood of

ARO transmission. Previous studies using MS2 to model the spread of *Clostridioides difficile* spores have also reported more frequent MS2 detections versus bacterial detections on HCP skin and clothing.^{19,35} However, in our study, both surrogate markers were present in all of the patient rooms, and only 7 rooms had surfaces positive for VRE, MRSA, or MSSA at baseline. Therefore, the lower rate of positive bacterial cultures was not unexpected.

In this study, both surrogate markers were identified less frequently among HCP caring for patients on contact precautions versus HCP caring for patients not on contact precautions, although the difference only reached significance for FP. We also observed that HCP caring for patients on contact precautions more frequently performed HH at room exit and that HCP who performed HH had fewer FP detections. Previous studies have also reported an association between contact precautions and HH

compliance^{36,37} and between HH and fewer MS2 detections on the hands of HCP.^{38,39} These findings suggest that both contact precautions and HH play an important role in preventing the spread of AROs, and they provide additional data to support the role of contact precautions in preventing ARO transmission.

Although we found no significant differences in the rate of FP detections among different types of HCP, MS2 was more frequently detected among physicians than among nurses. This observation may also be related to HH because nurses were more likely than physicians to perform HH at room exit. As in prior studies,⁴⁰ observed HCP HH compliance was low. However, differences in HH compliance by HCP job category suggest that a role exists for interventions promoting HH among all HCP.

A key strength of this study is the use of multiple surrogate markers and bacterial cultures, which helps to generate a more complete model of pathogen transmission. Other strengths include the real-world hospital setting and detailed HCP observations. This study also had several limitations. The small sample size may have limited the statistical power to detect differences in surrogate marker detections. This study also only included patients on contact precautions for VRE and MRSA, and we only tested for VRE, MRSA, and MSSA. Therefore, it is unclear how our findings would translate to other AROs, such as *C. difficile* and multidrug-resistant gram-negative bacteria. Finally, despite detailed HCP observations, it was not always possible to observe HH that occurred inside patient rooms when the door was closed. Therefore, we may have underestimated HCP HH compliance; however, our internal, routine HH observations indicate that HH compliance among hospital staff is less than ideal overall.

Despite these limitations, this study demonstrated transfer of both FP and MS2 beyond the initial areas of contamination inside patient rooms. Our findings suggest that both surrogate markers may be useful tools for studying ARO transmission. Larger studies using surrogate markers to assess ARO transmission and HCP cross-contamination are warranted, especially those focusing on the impact of contact precautions on ARO transmission.

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