

Contaminated Sinks May be an Environmental Source for Serial Transmission of Carbapenem-Resistant Enterobacteriaceae (CRE) to ICU Patients

iphe (**) **DUKE INFECTION PREVENTION** AND HOSPITAL EPIDEMIOLOGY

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Abstract

Background: We performed an investigation after noting an increase in hospital-onset (HO) KPC-producing Enterobacteriaceae (KPC-E) infections in patients admitted to a tertiary referral hospital in North Carolina.

Methods: We defined pre-outbreak (1/1/17-6/30/17), outbreak (7/1/17-10/31/17), and post-outbreak (11/1/17-7/31/18) phases. A clinical case was defined as any positive clinical culture for KPC-E. HO was defined as a positive clinical or surveillance culture collected on hospital day > 3Patients were mapped in space and time to inform targeted environmental sampling. Whole genome sequencing (WGS) was performed on selected KPC K. pneumoniae environmental and patient isolates to determine relatedness.

In October 2017, a CRE prevention bundle was implemented that included daily communication of CRE patient movement, increased audits/feedback of HCW compliance with hand hygiene, enhanced cleaning and disinfection in CRE rooms and high-risk units with bleach and UVC disinfection, and weekly rectal surveillance screens in 4 adult ICUs.

Results: 0.67 clinical cases of KPC-E per month were observed during the pre-outbreak period compared to 3.75 clinical cases of KPC-E per month during the outbreak period. Kp was the most common species (Figure 1). Mapping of patients revealed probable direct and indirect transmission between patients in multiple hospital units (Figure 2). 3 patients who were non-sequentially admitted to the same ICU room over a 12-week span acquired KPC Kp (Figure 2). Environmental cultures from the in-room sink drain and P-trap grew KPC Kp that was related to the patient isolates by WGS; the sink was removed. Although no additional clinical cases of KPC-E occurred after full implementation of the bundle and sink removal, we continued to observe acquisition of KPC-E rectal colonization in all 4 ICUs (Figure 3).

Conclusion: We describe a multi-species outbreak of KPC-E that was mitigated through evidence-based CRE control measures and removal of a colonized sink. However ongoing low-level presumed transmission of KPC points to persistent environmental sources. Additional study is needed to understand the prevalence of CRE in hospital sinks, factors that drive drain colonization, and contribution of CRE in a sink to nosocomial transmission.

Primary Objective To understand and mitigate transmission of KPCproducing Enterobacteriaceae (CPE) at a 957-bed tertiary care hospital using standard epidemiologic

Definitions

methods

- Outbreak periods
- Post-outbreak: 11/1/2017-7/31/2017 Clinical case: Any positive clinical culture for KPC-

CRE Pre

Daily communi Increased audi with hand hygi Enhanced clea patients with C bleach and UV Patient, family, hand hygiene Weekly rectal ICUs

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- Pre-outbreak: 1/1/2017-6/30/2017
- Outbreak: 7/1/2017-10/31/2017

producing *Enterobacteriaceae*

Rectal surveillance case: A positive rectal screening culture for KPC-producing *Enterobacteriaceae*

Hospital-onset (HO): Positive clinical or surveillance culture obtained on hospital day 3 or later

vention Bundle Elements	Date
ication of CRE patient movement	August 2017
its/feedback of HCW compliance ene and PPE compliance	August 2017
aning and disinfection in rooms of CRE and high-risk units using /-C light	August 2017
, and staff education regarding and PPE compliance	August 2017
surveillance screens in 4 adult	October 2017

Results

Pre-outbreak 0.67 clinical

Mapping of patients in time and space revealed probable direct and indirect transmission between patients in multiple hospital units (Figure 2)

I S. marcescens-Rectal

	3/2/2017		3/30/2017
Patient A Un	it 1		
Patient B Un	it 1		
Patient C Un	it 1		
Patient E Un	it 1		[
Patient F Un	it 1		* 3 pa
Patient G Un	it 2		same l
Patient H Un	it 2		onset
Patient J Un	it 2		
Patient K Un	it 3		
Pateint L Un	it 3		
Patient M Un	it 3		
Patient N Un	it 3		
Patient N Un	it 3		
Patient N Un	it 4		
Patient N Un	it 4		
Patient O Un	it 4		
Patient K Un	it 4		
Patient P Un	it 5		
Patient B Un	it 5		









Conclusions

- hospital transmission of CRE

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Targeted environmental sampling was performed in the room in Unit 2 with multiple patient acquisitions of KPC K pneumoniae and revealed KPCcolonization of in-room sink drain and P-trap of ICU room.

Additional point-prevalence sampling of in-room sinks in 3 units confirmed a high burden of in-room sink drain colonization.

Figure 3. Sub-clinical transmission of KPC-Producing Enterobacteriaceae in 4 Adult ICUs

We mitigated a multi-species outbreak of KPC-producing *Enterobacteriaceae* through implementation of evidence-based CRE control measures

Ongoing positive rectal surveillance screens demonstrate that sub-clinical transmission continues in high-risk units despite implementation of control measures

The high rate of in-room sink KPC colonization in 3 units implicates sinks as potential environmental reservoirs of CRE

Additional study is needed to understand the prevalence of CRE in hospital sinks, factors that drive drain colonization, and contribution of CRE in sink drains to

