

UK; ²John Farman Intensive Care Unit, Cambridge University Hospitals NHS Foundation Trust, Cambridge, UK; ³Department of Infectious Diseases, Cambridge University Hospitals NHS Foundation Trust, Cambridge, UK; ⁴Clinical Microbiology and Public Health Laboratory, Public Health England, Cambridge, UK; ⁵Department of Microbiology, Public Health England, Cambridge, UK; ⁶Wellcome Trust Sanger Institute, Cambridge, UK; ⁷London School of Hygiene and Tropical Medicine, London, UK

Session: 190. Resist! MDROs in Healthcare
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Background. The increasing incidence of carbapenemase-producing Enterobacteriaceae (CPE) is a global health concern, as treatment options are extremely limited. The prevalence of CPE in UK hospitals is unknown, as national screening guidelines only recommend screening in patients considered to be at high-risk of CPE. Patients in intensive care units (ICU) are at high-risk of healthcare-associated infections caused by multidrug-resistant organisms (MDRO).

Methods. We conducted a six-month prospective surveillance study to determine the prevalence of MDRO in a UK teaching hospital ICU. Between June and December 2016, all adult patients admitted to ICU were screened for MDRO on admission, on discharge, and weekly during their ICU stay. Surveillance samples included stool or rectal swabs, urine, sputum or tracheal aspirates, and wound swabs (if wounds were present). Isolates were characterized phenotypically before undergoing whole-genome sequencing (WGS), epidemiological, and phylogenetic analyses.

Results. During the first week of the study we identified stool carriage of a multidrug-resistant *Klebsiella pneumoniae* strain in two patients neither of whom had recognized risk factors for CPE. Both isolates were resistant to all antibiotics tested, apart from colistin, and were PCR-positive for the *bla*_{NDM-1} gene. Enhanced surveillance by the infection control team identified four additional patients in several wards who had stool carriage (*n* = 3) or bloodstream infection (*n* = 1) with a *bla*_{NDM-1} *K. pneumoniae* isolate. Epidemiological links were identified between these six patients. Five months later, a second outbreak of multidrug-resistant *K. pneumoniae* was detected, involving stool carriage by four patients on two different wards. Environmental screening identified environmental contamination with multidrug-resistant *K. pneumoniae* on one ward. DNA sequence analysis confirmed that a novel *bla*_{NDM-1} *K. pneumoniae* lineage (ST78) was responsible for both outbreaks in the hospital.

Conclusion. We identified two unsuspected *bla*_{NDM-1} *K. pneumoniae* outbreaks in patients with no recognized risk factors for CPE. This highlights the importance of prospective surveillance for MDRO in high-risk settings, such as ICUs, and supports the use of rapid WGS to support outbreak investigations in real-time.

Disclosures. All authors: No reported disclosures.

1698. Comparison of 30- and 90-Day Mortality Rates in Patients with Cultures Positive for Carbapenem-resistant Enterobacteriaceae and Acinetobacter in Atlanta, 2011–2015

Mary Elizabeth Sexton, MD^{1,2}, Chris Bower, MPH^{2,3,4}, Stephen Sukumaran, MPH^{2,3,4} and Jesse T. Jacob, MD^{1,2,5}; ¹Division of Infectious Diseases, Emory University School of Medicine, Atlanta, Georgia; ²Georgia Emerging Infections Program, Decatur, Georgia; ³Atlanta Veterans Affairs Medical Center, Decatur, Georgia; ⁴Atlanta Research and Education Foundation, Decatur, Georgia; ⁵Emory Antibiotic Resistance Center, Atlanta, Georgia

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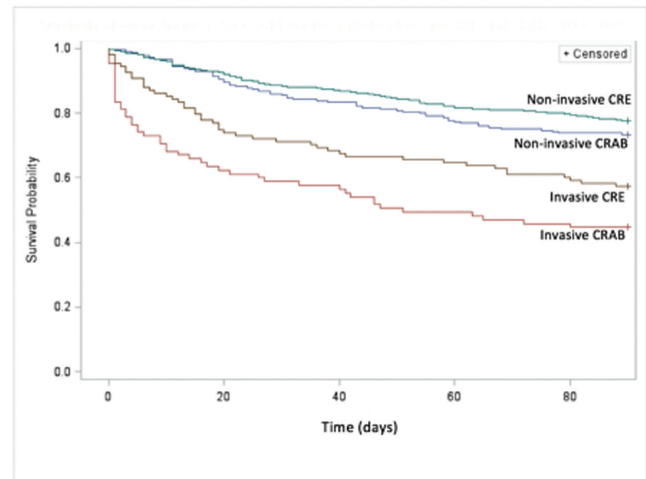
Background. Carbapenem-resistant Enterobacteriaceae (CRE) and *Acinetobacter baumannii* (CRAB) pose a threat to public health, but comparisons of disease burden are limited. We compared survival in patients following cultures positive for CRE or CRAB.

Methods. The Georgia Emerging Infections Program performs active population-based and laboratory-based surveillance for CRE and CRAB in metropolitan Atlanta, GA. Using standard CDC definitions, we included patients who had incident carbapenem-nonsusceptible *E. coli*, *Klebsiella* spp., *Enterobacter* spp., or *Acinetobacter baumannii* isolated from urine only (noninvasive infection) or a sterile site (invasive infection) between 8/2011 and 12/2015. Death dates, verified by Georgia Vital Statistics records, were used to calculate 30- and 90-day mortality rates. We used the chi-square test for mortality rates and the log-rank test for survival analysis to 90 days to compare patients with invasive CRAB, noninvasive CRAB, invasive CRE, and noninvasive CRE.

Results. There were 535 patients with CRE (87 invasive, 448 noninvasive) and 279 (78 invasive, 201 noninvasive) with CRAB. Nearly all patients with CRE and CRAB had healthcare exposures (97.2% vs. 100%) and most were immunosuppressed (62.6% vs. 56.3%). Both 30-day (24.4% vs. 18.3%, *p* = 0.04) and 90-day (37.6% vs. 30.5%, *p* = 0.04) mortality were higher in patients with CRAB than CRE. Patients with invasive infections were more likely to die at 90 days than those with noninvasive infections (53.3% vs. 38.4%, *p* < 0.0001). Overall mortality rates for invasive infection were similar between CRAB and CRE at 30 (44.9% vs. 34.5% *p* = 0.2) and 90 days (59.0% vs. 48.3%, *p* = 0.2). Using survival analysis at 90 days, invasive CRAB had the worst outcomes, followed by invasive CRE, noninvasive CRAB and noninvasive CRE (*p* < 0.0001, see Figure).

Conclusion. Ninety-day mortality for invasive infections with CRE and CRAB was ~50%, and patients with CRAB had lower survival than those with CRE, suggesting that prevention efforts may need to prioritize CRAB as highly as CRE in facilities with endemic CRAB. With the high proportion of healthcare exposures and immunosuppression, these infections may signify poor prognosis or directly contribute to mortality.

Figure. Survival Following Incident Cases of Invasive and Non-Invasive Carbapenem-resistant Enterobacteriaceae (CRE) and *Acinetobacter baumannii* (CRAB), Atlanta, 2011-2015



Disclosures. All authors: No reported disclosures.

1699. Prevalence and Acquisition of MRSA During Incarceration at a Large Inner-city Jail

Kyle J Popovich, MD MS FIDSA^{1,2}, Chad Zawitz, MD², Alla Aroutcheva, MD, PhD^{1,2}, Darjai Payne, BS, MPH¹, Michael Schoeny, PhD¹, Emma Richardson, BS¹, Mary K. Hayden, MD, FIDSA, FSHEA¹, Bala Hota, MD, MPH^{1,2}, Robert A. Weinstein, MD, FIDSA, FSHEA, FACP^{1,2}, ¹Rush University Medical Center, Chicago, Illinois; ²Stroger Hospital of Cook County, Chicago, Illinois

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Background. USA300 MRSA is endemic in certain communities, with congregate settings such as urban jails potentially facilitating spread. The extent of MRSA transmission in jail is unclear, a controversy that impacts prevention strategies. We determined the prevalence of MRSA colonization at jail entrance and defined the acquisition rate during incarceration.

Methods. Men incarcerated at the Cook County Jail, one of the largest US single-site jails, were enrolled within 72 hours of intake. Surveillance cultures (nares, throat, groin) were collected to determine prevalence of MRSA colonization. A survey was administered to identify predictors of colonization. Detainees still in jail at Day30 had cultures repeated to determine MRSA acquisition rate. Univariate and multivariate analysis was performed to identify predictors of MRSA colonization.

Results. A total of 402 men (447 unique incarcerations) have so far been enrolled (77% AA, 11% Hispanic) with 92% previously in jail (20% in past 6 months). The prevalence of MRSA colonization at intake was 18.6% (83/447), with 39% of those colonized solely in the throat or groin. At 30 days: 10% (9/92) of initially negative men acquired MRSA; 14 admission positives remained colonized while 11 lost colonization. On univariate (Table), predictors of MRSA colonization at entrance to the jail were: methamphetamine use (METH), unstable housing, current skin infection, and care at an outpatient Clinic A that emphasizes comprehensive care to the LGBTQ community. In this cohort, METH use was associated with reporting being a man who has sex with men vs. not (35% vs. 9%, *P* < 0.001) and was common among men with care at Clinic A (18% vs. 3%, *P* < 0.001). On multivariate with adjustment for race/ethnicity and HIV status, current skin infection and care at clinic A were associated with MRSA. Preliminarily, sharing personal items was associated with MRSA acquisition at Day30 (OR = 5.6, 95% CI, 1.3, 23.3, *P* = 0.02).

Conclusion. We found that a relatively high proportion of individuals enter the jail colonized with MRSA and the jail may amplify rates. Entrance colonization risk factors point to possible community reservoirs. Enrollment is ongoing but results suggest an intervention in jail could impact MRSA rates in the jail and in the surrounding community.