ID WEEK 2018

ORAL ABSTRACT

109. Differences in Gram-Negative Antibiotic Susceptibility Among Patients Receiving Fecal Microbiota Transplant for *Clostridioides difficile* <u>Michael Woodworth</u>, MD, MSc¹; Tiffany Wang, MD²; Divyanshu Raheja, MPH²; Alex Waldman, BS²; Rachel Friedman-Moraco, MD¹; Allen Graham, BA³; Tanvi Dhere, MD⁴ and Colleen Kraft, MD, MSc^{1,3}, 'Medicine, Division of Infectious Disease, Emory

University School of Medicine, Atlanta, Georgia, ²Emory University School of Medicine, Atlanta, Georgia, ³Pathology, Emory University School of Medicine, Atlanta, Georgia, ⁴Division of Digestive Diseases, Emory University School of Medicine, Atlanta, Georgia

Session: 30. Healthcare Epidemiology: Hot Topics

Thursday, October 4, 2018: 8:45 AM

Background. Decreases in multidrug-resistant organism (MDRO) colonization and antibiotic resistance gene abundance have been reported after fecal microbiota transplantation (FMT), but data on clinical microbiology culture and susceptibility results after FMT are limited.

Methods. We retrospectively reviewed the available microbiology results for patients who underwent FMT for recurrent *Clostridioides difficile* infection (RCDI) at Emory University from July 7, 2012 until December 2017 and had microbiology results within 1 year pre- and post-FMT. Demographic and clinical characteristics were abstracted by trained reviewers, and statistical tests of differences in central tendency were tested with Wilcoxon signed-rank tests.

Results. Of 236 unique patients undergoing FMT during the study period, 18 had growth of Gram-negative bacteria on culture pre- and post-FMT. Of these, 8 had Gram-negative growth in urine culture (the most common site) pre- and post-FMT. Fourteen (14/18, 78%) patients were female, 4/18 (22%) were black, 14/22 (78%) were white, and 18/18 (100%) were non-Hispanic. The mean number of CDI episodes prior to first FMT was 4 (range 3–7 episodes). Differences in counts of susceptible, intermediate, and resistant susceptibility test results before and after FMT are shown in Figures 1 and 2. Although a trend in reduction of resistant reports is visually suggested, this was not statistically significant by Wilcoxon signed-rank testing (P = 0.10 for all cultures, P = 0.21 for urine). Ten patients had pre-FMT micro results and no micro results after FMT, but reduction of count of infectious syndromes in FMT could not be tested with this study design. Abstraction of viral quantitative PCR results did not suggest clinical recognition of new infection or reactivation of viruses after FMT.

Conclusion. FMT may reduce clinical burden of antimicrobial resistance, but statistically significant differences in resistance were not detected in this study. Further study with RCTs is needed.



Figure 1: Gram-negative culture results from all sites within 1-year pre- and post-FMT, 2012–2017.



Figure 2: Gram-negative urine culture results within 1-year pre- and post-FMT, 2012–2017.

Disclosures. All authors: No reported disclosures.

Open Forum Infectious Diseases[®] 2018;1(S1):S1–75

© The Author(s) 2018. Published by Oxford University Press on behalf of Infectious Diseases Society of America. This is an Open Access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs licence (http://creativecommons.org/licenses/ by-nc-nd/4.0/), which permits non-commercial reproduction and distribution of the work, in any medium, provided the original work is not altered or transformed in any way, and that the work is properly cited. For commercial re-use, please contact journals.permissions@oup.com D0I: 10.1093/ofd/ofx209

110. The Burden and Preventability of Sepsis-Associated Mortality in 6 US Acute Care Hospitals

Chanu Rhee, MD, MPH¹; Travis Jones, PharmD²; Yasir Hamad, MD³; Anupam Pande, MD, MPH⁴; Jack Varon, MD⁵; Cara O'Brien, MD⁶; Deverick J. Anderson, MD, MPH, FIDSA, FSHEA⁷; David K. Warren, MD, MPH, FIDSA, FSHEA⁸, Raymund Dantes, MD, MPH⁹, Lauren Epstein, MD, MSc¹⁰ Michael Klompas, MD, MPH, FRCPC, FIDSA¹, ¹Department of Population and Medicine, Harvard Medical School and Harvard Pilgrim Health Care Institute, Boston, Massachusetts, ²Duke Antimicrobial Stewardship Outreach Network, Duke University Medical Center, Durham, North Carolina, ³Department of Medicine, Washington University School of Medicine, St. Louis, Missouri, ⁴Infectious Diseases, Washington University School of Medicine, St. Louis, Missouri, ⁵Department of Medicine, Brigham and Women's Hospital, Boston, Massachusetts, 6Department of Medicine, Duke University School of Medicine, Durham, North Carolina, 7Division of Infectious Diseases, Duke University School of Medicine, Durham, North Carolina, ⁸Division of Infectious Diseases, Washington University School of Medicine, St. Louis, Missouri, 9Department of Medicine, Emory University School of Medicine, Atlanta, Georgia, ¹⁰Division of Healthcare Quality Promotion, Centers for Disease Control and Prevention, Atlanta, Georgia

Session: 30. Healthcare Epidemiology: Hot Topics Thursday, October 4, 2018: 8:45 AM

Background. Sepsis is considered a leading cause of preventable death, but the actual burden of sepsis mortality is difficult to measure using administrative data or death certificates. We analyzed the prevalence, underlying causes, and preventability of deaths due to sepsis in acute care hospitals using detailed medical record reviews.

Methods. We randomly selected 577 adult patients who died in-hospital or were discharged to hospice in 2014–2015 at 6 US academic and community hospitals for medical record review. Cases were reviewed by experienced clinicians for sepsis during hospitalization (using Sepsis-3 criteria), terminal conditions on admission (defined using hospice-qualifying criteria), immediate and underlying causes of death, and suboptimal sepsis care (delays in antibiotics, inappropriate antibiotic therapy, inadequate source control, or other medical errors). The overall preventability of death was rated on a 6-point Likert scale (from definitely not preventable to definitely preventable) taking into account comorbidities, severity of illness, and quality of care.

Results. Sepsis was present in 302/577 (52%) hospitalizations ending in death or discharge to hospice and was the immediate cause of death in 199 cases (35%) (Figure 1A). Underlying causes of death in sepsis patients included solid cancer (21%) and chronic heart disease (15%), and hematologic cancer (10%) (Figure 1B). The median age of sepsis patients who died was 73 (IQR 62–84). Terminal conditions were present in 122/302 (40%) sepsis deaths, most commonly end-stage cancer (26% of cases). Suboptimal care was identified in 68 (23%) of sepsis deaths, most commonly delays in antibiotics (11% of cases). However, only 4% of sepsis deaths were definitely or likely preventable and an additional 8% were considered possibly preventable with optimal clinical care (Figures 2 and 3).

Conclusion. Our findings affirm that sepsis is the most common cause of death in hospitalized patients. Most patients that died with sepsis were elderly with severe comorbidities, but up to 1 in 8 sepsis deaths were felt to be potentially preventable with better hospital-based care. These findings may inform resource allocation and expectations surrounding the impact of hospital-based sepsis treatment initiatives.

Figure 1. Immediate and Underlying Causes of Death

A. IMMEDIATE CAUSE OF DEATH (ALL PATIENTS, N=577)

