



Thinking Outside The Bowel: *Clostridium difficile* Bacteremia Case Series

Adam Pettigrew¹, Ripal Jariwala², Kristen Zeitler², Jose Montero¹, Sandra Gompf^{1,3}, John Toney^{1,3}

¹Division of Infectious Diseases & International Medicine, University of South Florida;

²Department of Pharmacy, Tampa General Hospital; ³Infectious Disease Section, James A. Haley Veterans' Hospital

Background

- *Clostridium difficile* gastrointestinal infection (CDI) is the most common hospital-acquired infectious disease, yet *Clostridium difficile* bacteremia (CDB) is exceedingly rare
- CDB risk factors, mortality rate, and modalities of treatment are not well defined
- CDB may occur in individuals with significant co-morbidities, including gastrointestinal pathologic conditions^{1,2}
- Antimicrobial therapy can be variable and adapted for polymicrobial bacteremia²
- Outcomes may be impacted by early diagnosis as well as severity of underlying conditions^{1,2}

Methods

- Retrospective, IRB-approved, chart review
- Adult patients with diagnosis of CDB admitted to Tampa General Hospital or James A. Haley VA Hospital from 2011 through 2017
- Catalogued variables:
 - Patient age & co-morbid conditions
 - Gastrointestinal (GI) disruption (perforated viscous, GI bleeding, abdominal malignancy)
 - Diarrhea at time of CDB
 - Proton Pump Inhibitor (PPI) use
 - Previous antibiotics
 - Prior history of CDI
 - Treatment courses & outcomes

Conclusions

- CDI is the most common cause of hospital acquired infection, though rarely causes bacteremia
- Notable findings in our population included older age, concomitant malignancy, evidence of GI disruption, and prior exposure to PPIs and antibiotics
- Antibiotics chosen to treat CDB were IV metronidazole and/or IV vancomycin, with other broad spectrum antibiotics utilized due to polymicrobial bacteremia
- CDB is associated with a high mortality rate and is commonly manifested as a polymicrobial bloodstream infection
- This is one of the larger case series that adds to the scant literature characterizing patients diagnosed with CDB

References

1. Mattila E, Arkkila P, Mattila PS, et al. Extra-intestinal *Clostridium difficile* infections. *Clin Infect Dis* 2013; 57 (6): e148-53.
 2. Doufair M, Eckert C, Drieux L, et al. *Clostridium difficile* bacteremia: Report of two cases in French hospitals and comprehensive review of the literature. *ID Cases* 2017; 8: 54-62.

Results

| Patient Age | Pertinent Co-Morbid Conditions | GI Disruption | Diarrhea at time of CDB | Prior PPI | Abx Exposure in Previous 30 Days | Prior Positive CDI by Testing | Current CDI Testing During Episode | Other Microbes in Bloodstream | Management of Bacteremia | Outcome |
|-------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------|-----------|----------------------------------|-------------------------------|------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|----------|
| 70 | <ul style="list-style-type: none"> • Diabetes mellitus (DM) with foot infection • Peripheral vascular disease (PVD) | No | No | Yes | No | Yes | No | <ul style="list-style-type: none"> • <i>Clostridium tertium</i> • Diphtheroids | Prior broad antibiotics de-escalated to metronidazole x 2 weeks BKA performed | Survived |
| 81 | <ul style="list-style-type: none"> • DM with chronic kidney disease • PVD • Coronary artery disease (CAD) | <ul style="list-style-type: none"> • Pneumoperitoneum • Necrotic ileum on ex-lap | No | Yes | No | No | Yes (+ PCR) | <ul style="list-style-type: none"> • <i>Enterococcus faecalis</i> • <i>Klebsiella pneumoniae</i> • Coagulase negative staphylococcus | Doripenem x 7 days, Metronidazole x 28 days, PO vancomycin x 14 days | Survived |
| 35 | <ul style="list-style-type: none"> • Testicular cancer 2 years prior • CAD • PTSD (foreign body ingestions) | <ul style="list-style-type: none"> • Abdominal pain with recent GI bleed • Esophagitis, gastritis, and duodenitis by EGD | Yes | Yes | Yes | No | No | <i>Candida dublinensis</i> | Metronidazole x 2 weeks, Fluconazole x 2 weeks | Survived |
| 63 | <ul style="list-style-type: none"> • Endometrial cancer • Bilateral hydronephrosis • Pulmonary embolism | <ul style="list-style-type: none"> • TAH/BSO with colorectal anastomosis • Takeback for appendectomy, L colectomy with colostomy | No | Yes | Yes | No | Yes (- PCR) | <ul style="list-style-type: none"> • <i>Enterococcus faecalis</i> • <i>Eubacterium lentum</i> | IV Vancomycin, Ciprofloxacin + Metronidazole x 10 days | Survived |
| 58 | <ul style="list-style-type: none"> • Uterine cancer s/p TAH/BSO • Receipt of chemotherapy & radiation with progression of disease | Metastasis to peritoneum and mesentery | Yes | No | No | Yes | Yes (+ PCR) | <ul style="list-style-type: none"> • <i>Clostridium cadavaris</i> • <i>Clostridium septicum</i> x2 • <i>Clostridium butyricum</i> • <i>Clostridium innocuum</i> • <i>Clostridium perfringes</i> • <i>Bacteroides vulgatus</i> | Doripenem + IV Vancomycin, then switched to IV Metronidazole until CMO | Died |
| 65 | <ul style="list-style-type: none"> • Metastatic colon cancer with hepatic wedge resection • DM • CAD | <ul style="list-style-type: none"> • Colon cancer • Surgery (hepatic wedge resection) | No | Yes | Yes | Yes | Yes (+ PCR; - toxin) | None | IV Vancomycin + Piperacillin / tazobactam, then Tigecycline until CMO | Died |
| 74 | <ul style="list-style-type: none"> • Cervical cancer s/p hysterectomy and pelvic radiation • CAD • Deep vein thrombosis | Colo-vesicular fistula | Yes | Yes | Yes | No | No | None | Linezolid switched to IV Metronidazole + Fluconazole until CMO | Died |