Fatal Spontaneous Clostridium bifermantans Necrotizing Endometritis: A Case Report and Literature Review of the Pathogen

Citation

Published Version
doi:10.1093/ofid/ofw095

Permanent link
http://nrs.harvard.edu/urn-3:HUL.InstRepos:27822138

Terms of Use
This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA

Share Your Story
The Harvard community has made this article openly available. Please share how this access benefits you. Submit a story.

Accessibility
Fatal Spontaneous Clostridium bifermentans Necrotizing Endometritis: A Case Report and Literature Review of the Pathogen

Andrew Hale,1 James E. Kirby,2,3 and Mary Albrecht1,4

1Division of Infection Diseases, 2Department of Clinical Microbiology, Beth Israel Deaconess Medical Center, Departments of 3Pathology, and 4Medicine, Harvard Medical School, Boston, Massachusetts

Clostridium bifermentans is a rare pathogen in humans. A fatal case of fulminating endometritis with toxic shock and capillary leak secondary to C bifermentans infection in a young woman is described, and this is compared to all 13 previously described cases of C bifermentans infection.

Keywords. capillary leak; Clostridium bifermentans; Clostridium sordellii; hyperleukocytosis; toxic shock syndrome.

In October 2014, a 33-year-old woman with history of abnormal uterine bleeding status post an uncomplicated endometrial ablation 5 months prior developed new onset dysuria and grayish, malodorous vaginal discharge. Her white blood cell count was 16 × 10^3 cells/mm^3 and her hematocrit was 39%. She was prescribed levofloxacin but never filled it. Three days later, she was hospitalized after multiple syncopal events at home. On admission, blood pressure was 74/24 mmHg. She was afebrile, without rash, and extremities were cool. Abdominal exam was notable for obesity but was otherwise benign. Pelvic examination showed no cervical motion tenderness, cervical discharge, or other abnormality. Urine β-human chorionic gonadotropin was negative. White blood cell count was 60 × 10^3 cells/mm^3; hematocrit 42%; platelets 38 × 10^3 cells/mm^3; and lactate 6.5 mmol/L. Computed tomography (CT) scan of the abdomen and pelvis showed ascites, and a pelvic ultrasound was unremarkable. Empiric antibiotic therapy with vancomycin, cefepime, and metronidazole was initiated, and she was transferred to the intensive care unit. Twelve hours after admission, her white blood cell count rose to 113 × 10^3 cells/mm^3, and hematocrit was 58%, despite having received 12 liters of intravenous (IV) normal saline. Lactate increased to 13.5 mmol/L. Tobramycin, clindamycin, and doxycycline were added for sepsis. Despite vasopressor support, she required massive IV fluid resuscitation to maintain pressures and perfusion, receiving 26 liters during the first 24 hours and 51 liters by 72 hours. In addition, she showed signs of disseminated intravascular coagulation (DIC) and received 46 units of blood product (fresh frozen plasma, platelets, and cryoprecipitate) over 72 hours. Transthoracic echocardiogram showed a large pericardial effusion with cardiac tamponade, which required pericardial drain placement with removal of 1500 mL transudative fluid. She also developed large bilateral pleural effusions and required bilateral chest tube placement with 4 liters per day transudative fluid output. Given the profound intra-abdominal fluid, there was concern for potential abdominal catastrophe; thus, she had an exploratory laparotomy, which was negative for any signs of intraperitoneal infection or viscous perforation, although there was massive ascites. The uterus was hyperemic but not grossly infected or boggy. Vaginal swabs did not isolate Staphylococcus aureus or yeast; cervical swabs were negative for gonorrhea and chlamydia by nucleic acid amplification testing. Blood smear showed marked neutrophilia and no evidence of hematologic malignancy. Bone marrow biopsy was negative for malignancy. Human immunodeficiency virus-1 antibody was negative. Blood, urine, pleural fluid, pericardial fluid, peritoneal fluid, and sputum cultures were negative for microorganisms. She developed profound anasarca, and a repeat pelvic or ophthal-mologic exam was unable to be performed because of massive edema. Her clinical picture was felt to be consistent with a toxin-mediated process, potentially related to Clostridium sordellii given how this pathogen typically presents. Thus, to remove potential toxin, empiric plasmapheresis was started on hospital day 3, which showed immediate improvement in hemodynamics, and fluid was able to be removed with continuous venovenous hemofiltration. On hospital day 4, her edema had improved and the team was able to examine her pupils, which were found to be fixed and dilated. A CT head scan showed diffuse cerebral edema and tonsillar herniation. A family meeting was convened, life support was withdrawn, and she died shortly thereafter. An autopsy demonstrated diffuse edema and evidence of DIC in all organs. Diffuse endometrial necrosis was noted on histopathological examination of the uterus. Special stains demonstrated large, boxcar-shaped, Gram-positive rods within endometrial tissue. Uterine tissue submitted to the Centers for Disease Control and Prevention confirmed the presence of clostridial species by immunohistochemical

Received 16 December 2015; accepted 9 May 2016.

Correspondence: A. Hale, Lowry Medical Office Building, 110 Francis Street, Suite GB, Boston, MA 02215 (ahale@bidmc.harvard.edu).

Open Forum Infectious Diseases®
© The Author 2016. Published by Oxford University Press on behalf of the 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. DOI: 10.1093/ofid/ofw095
staining within the areas of endometrial necrosis (Figure 1). Although polymerase chain reaction (PCR) assay specific for C. sordellii was negative, wide-range 16 S PCR assay of endometrial tissue was positive for Clostridium bifermentans. Postmortem endometrial cultures were negative.

**DISCUSSION**

The Clostridium family represents a diverse group of Gram-positive, spore-forming, obligate anaerobic bacteria that are found widely throughout the environment and are known to secrete a wide array of toxins [1]. Clostridium bifermentans was first isolated in 1902 in putrefied butcher’s meat [2]. It is found in sewage, soil, and occasionally the intestinal flora of humans. The frequency of C. bifermentans causing human infection is quite rare; our search of PubMed revealed only 13 prior case reports. Sites of infection are diverse (Table 1) [2–14]. Of these prior reported cases, only 1 infection, presenting with necrotizing pneumonia and empyema, was fatal [5] (7.7% case fatality rate). A disproportionate number of infections were in men (85%; see Table 1). The case we now report is quite unique. Histopathology showed infection of the uterus, a site of infection not previously described. In addition, our patient’s course was fulminant and rapidly fatal, which is not a characteristic feature in prior cases of C. bifermentans infection. Our patient’s course was much more in line with what has previously been described in C. sordellii infection, which is well known to cause a fulminant endometritis, typically in young women after medical abortion [15], although it can occur spontaneously [16]. Our patient had an endometrial ablation 5 months before her presentation, which was felt to be too distant in time to be a definitive risk factor. Clostridium sordellii is known to cause a severe leukocytosis in part related to production of the neuraminidase NanS, which stimulates promyelocytic proliferation and prevents margination and movement of leukocytes out of the intravascular space [17]. Clostridium sordellii also elaborates lethal toxin, which undermines the actin cytoskeleton at the cellular level, and is believed through this activity to compromise endothelial barrier integrity [18]. Lethal toxin exhibits a marked propensity for inducing rapid morbidity and contributes to the profound capillary leak, hemoconcentration, and toxic shock syndrome often seen in C. sordellii infections [19]. The constellation of findings in our patient (severe capillary leak, profound leukocytosis and hemoconcentration, and improvement with plasmapheresis) was very suspicious for C. sordellii. However, unexpectedly, PCR testing revealed evidence of C. bifermentans in endometrial tissue. Clostridium sordellii and Clostridium septicum-specific PCR assays were negative. We thus surmise that this particular strain of C. bifermentans may have elaborated toxins similar to the C. sordellii lethal toxin and NanS, although this conjecture remains unproven. Of note, however, genetic exchange between large toxin protein producing strains of Clostridium has been previously suggested [20]. Moreover, within

---

**Table 1. Review of 13 Prior Cases of Clostridium bifermentans Infection**

<table>
<thead>
<tr>
<th>Author Name</th>
<th>Publication Year</th>
<th>Site of Infection</th>
<th>Patient Age in Years</th>
<th>Patient Sex</th>
<th>Maximum WBC in $10^3$ Cells/mm$^3$</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bitner</td>
<td>1971</td>
<td>Abdominal wall</td>
<td>45</td>
<td>Female</td>
<td>19</td>
<td>Survived</td>
</tr>
<tr>
<td>Nolan</td>
<td>1972</td>
<td>Septic arthritis of knee</td>
<td>18</td>
<td>Male</td>
<td>10</td>
<td>Survived</td>
</tr>
<tr>
<td>Misra</td>
<td>1980</td>
<td>Necrotizing pneumonia and empyema</td>
<td>41</td>
<td>Female</td>
<td>52</td>
<td>Death</td>
</tr>
<tr>
<td>Panwalker</td>
<td>1983</td>
<td>Necrotizing pneumonia and empyema</td>
<td>60</td>
<td>Male</td>
<td>24</td>
<td>Survived</td>
</tr>
<tr>
<td>Pencek</td>
<td>1986</td>
<td>Brain abscess</td>
<td>36</td>
<td>Male</td>
<td>9</td>
<td>Survived</td>
</tr>
<tr>
<td>Kolander</td>
<td>1989</td>
<td>Endocarditis</td>
<td>28</td>
<td>Male</td>
<td>7</td>
<td>Survived</td>
</tr>
<tr>
<td>Nachman</td>
<td>1989</td>
<td>Liver abscess</td>
<td>6</td>
<td>Male</td>
<td>25</td>
<td>Survived</td>
</tr>
<tr>
<td>Rechtman</td>
<td>1991</td>
<td>Abdominal abscess</td>
<td>65</td>
<td>Male</td>
<td>9</td>
<td>Survived</td>
</tr>
<tr>
<td>Moyano</td>
<td>1994</td>
<td>Endocarditis</td>
<td>26</td>
<td>Male</td>
<td>16</td>
<td>Survived</td>
</tr>
<tr>
<td>Rehany</td>
<td>1994</td>
<td>Panophthalmitis</td>
<td>13</td>
<td>Male</td>
<td>9</td>
<td>Survived</td>
</tr>
<tr>
<td>Scanlan</td>
<td>1994</td>
<td>Osteomyelitis and bacteremia</td>
<td>81</td>
<td>Male</td>
<td>3</td>
<td>Survived</td>
</tr>
<tr>
<td>Chaudry</td>
<td>2014</td>
<td>Endocarditis</td>
<td>22</td>
<td>Male</td>
<td>7</td>
<td>Survived</td>
</tr>
<tr>
<td>Edagiz</td>
<td>2015</td>
<td>Empyema</td>
<td>60</td>
<td>Male</td>
<td>18</td>
<td>Survived</td>
</tr>
</tbody>
</table>

Abbreviation: WBC, white blood cells.
the Clostridium phylogeny, C bifermentans and C sordellii are closely related and were not identified as separate species until 1962 [21]. It is also remarkable that in the one prior fatal case of C bifermentans, the white blood cell count was 52 × 10^3 cells/mm^3; no other reported cases until ours showed an extreme leukocytosis. This may suggest that the toxins alluded to above may be present in only a small, lethal subset of strains, although that is also unproven. Because no cultures were positive in our case, assays to study toxin elaboration were not possible.

CONCLUSIONS

Clostridium bifermentans is a rare cause of infection in humans. Our case represents a novel manifestation of C bifermentans in regards to both site and severity. Further characterization of this rare pathogen is warranted.

Acknowledgments

Potential conflicts of interest. All authors: No reported conflicts. All authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest.

References