Bacillus subtilis variant natto Bacteremia of Gastrointestinal Origin, Japan

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We report a case of bacteremia caused by Bacillus subtilis variant natto after a gastrointestinal perforation in a patient in Japan. Genotypic and phenotypic studies of biotin identified B. subtilis var. natto. This case and 3 others in Japan may have been caused by consuming natto (fermented soybeans).

Bacillus subtilis is a gram-positive, rod-shaped, spore-forming bacterium temporarily present in the human gastrointestinal tract (1). The presence of B. subtilis in clinical specimens indicates contamination, but rare cases of bacteremia have been reported in Japan (2). Previous reports have attributed bacteremia in Japan to gastrointestinal origin but of unknown cause. We identified a case of B. subtilis variant natto bacteremia in a patient in Japan.

In May 2021, a 56-year-old woman was referred to the Japanese Red Cross Musashino Hospital (Musashino-shi, Tokyo, Japan) for a 2-day history of abdominal pain after having taken barium for gastric radiographic examination. The patient had a history of hypertension and ate natto (fermented soybeans) almost every day. At admission, the patient exhibited spontaneous abdominal pain, muscular defense, and rebound tenderness. Laboratory findings showed a decreased leukocyte count (1,800 cells/µL, reference range 3,300–8,600 cells/µL) and mildly increased C-reactive protein concentration (0.75 mg/dL, reference range 0–0.14 mg/dL). Contrast-enhanced computed tomography revealed contrast accumulation in the colon and free air around the sigmoid rectum. Lower gastrointestinal perforation and generalized peritonitis were suspected, and 2 sets of blood cultures were obtained. Emergency proctosigmoidectomy (Hartmann surgery) was performed on the same day, and perforation of the sigmoid colon was confirmed.

Intravenous antimicrobial treatment was initiated. Initial treatment was piperacillin/tazobactam (18 g/d). On day 5, because both blood culture sets were positive for gram-positive rod bacteria, teicoplanin (800 mg/d) was added. On day 11, only B. subtilis was isolated from the culture by matrix-assisted laser desorption/ionization-time of flight mass spectrometry, and the antimicrobial drugs

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**Figure.** Bacillus subtilis cultures on E9 minimal medium agar plates with and without biotin. From left to right in each column, 0.5 McFarland standard was diluted ×1, ×10, and ×10², and 10 µL was incubated at 35°C for 72 hours under aerobic conditions. The isolate showed a biotin requirement. Isolate, Bacillus subtilis variant natto from patient in Japan with bacteremia of gastrointestinal origin; natto, B. subtilis var. natto standard strain; subtilis, B. subtilis subspecies subtilis standard strain.
were changed to ampicillin/sulbactam (12 g/d) as indicated by antimicrobial susceptibility testing by broth microdilution (Appendix Table, https://wwwnc.cdc.gov/EID/article/28/8/21-1567-App1.pdf). B. subtilis was also detected along with multiple other bacteria by culture of ascites fluid collected intraoperatively. After 39 days of antimicrobial therapy, the patient was discharged.

We investigated whether the blood culture isolate was B. subtilis var. natto. DNA analysis showed that in the bioF region, the isolate was 100% homologous to the B. subtilis var. natto standard strain. Compared with the B. subtilis subspecies subtilis standard strain, the isolate had ≈50 fewer bases and the bioV region of the isolate had a single-nucleotide mutation that resulted in a termination codon for amino acid synthesis (Appendix Figures 1–4). The isolate and B. subtilis var. natto standard strain grew abundantly on a biotin-supplemented medium but did not thrive on a nonsupplemented medium (Figure).

Our biotin gene and biotin requirement testing confirmed that the isolate in this case was B. subtilis var. natto. Previous genotypic and phenotypic studies on biotin were helpful in identifying this variant. Kubo et al. reported that natto-fermented B. subtilis requires biotin and that nonfermented B. subtilis does not (3). bioF and bioV are biotin biosynthetic operons in B. subtilis (4). Compared with the B. subtilis subsp. subtilis standard strain, the 2 biotin genes of the isolate in this study and the B. subtilis var. natto standard strain were partially defective. According to the biotin requirement test, the isolate required biotin.

We conclude that this case of bacteremia caused by B. subtilis var. natto resulted from a gastrointestinal perforation. In Japan, the most common causative organism of community-acquired bloodstream infections is gram-negative Escherichia coli (25.4%); gram-positive bacilli rarely induce bacteremia (2.7%) (5). B. subtilis bacteremia typically originates from the gastrointestinal tract (2); Tamura et al. have reported 3 cases of B. subtilis bacteremia arising from the gastrointestinal tract (6). In patients with gastrointestinal bacteremia, the causative organism differs according to the food consumed (7). Oggoni et al. reported a case of B. subtilis bacteremia caused by probiotics (8). However, the patient that we report was not taking any probiotics but frequently ate natto. Most of the previously reported cases of B. subtilis bacteremia in Japan (2,6) were possibly related to natto consumption, although dietary history was not mentioned in their reports.

This case of bacteremia caused by B. subtilis var. natto resulted from gastrointestinal tract perforation. Genotypic and phenotypic studies on biotin effectively identified B. subtilis var. natto. In Japan, natto consumption is common, and B. subtilis bacteremia of gastrointestinal origin is most likely associated with B. subtilis var. natto.

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Mr. Tanaka is a pharmacist at the Japanese Red Cross Musashino Hospital in Musashino-shi, Tokyo, Japan. His main research interest is microbiology.

References


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Appendix

Appendix Table. Minimum inhibitory concentrations of each antimicrobial agent and their interpretation against the Bacillus subtilis var. natto strains isolated from the patient*

<table>
<thead>
<tr>
<th>Antimicrobial agent</th>
<th>MIC (µg/mL)</th>
<th>Interpretation†</th>
</tr>
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<tbody>
<tr>
<td>Penicillin G</td>
<td>≤0.12</td>
<td>S</td>
</tr>
<tr>
<td>Ampicillin</td>
<td>≤0.06</td>
<td>S</td>
</tr>
<tr>
<td>Imipenem</td>
<td>≤1</td>
<td>S</td>
</tr>
<tr>
<td>Gentamicin</td>
<td>≤1</td>
<td>S</td>
</tr>
<tr>
<td>Erythromycin</td>
<td>≤0.25</td>
<td>S</td>
</tr>
<tr>
<td>Clindamycin</td>
<td>≤0.25</td>
<td>S</td>
</tr>
<tr>
<td>Levofloxacin</td>
<td>≤0.5</td>
<td>S</td>
</tr>
</tbody>
</table>

*MIC, minimum inhibitory concentration; S, susceptible; R, resistant.
†Interpretation (susceptible, intermediate, or resistant) was based on the Clinical and Laboratory Standards Institute criteria (M45).

Appendix Figure 1. The isolate showed 100% homology to Bacillus subtilis var. natto standard strain in the bioF sequence. natto, Bacillus subtilis var. natto standard strain.
Appendix Figure 2. The isolate showed a 50-nt deletion in the bioF sequence compared to *Bacillus subtilis* subsp. *subtilis* standard strain. subtilis, *Bacillus subtilis* subsp. *subtilis* standard strain.

Appendix Figure 3. The isolate showed 100% homology to *Bacillus subtilis* var. *natto* standard strain in the bioW sequence. natto, *Bacillus subtilis* var. *natto* standard strain.
Appendix Figure 4. The isolate showed a single nucleotide mutation that resulted in a termination codon and stopped amino acid synthesis compared to *B. subtilis* subsp. *subtilis* standard strain in the *bioW* sequence. *subtilis*, *Bacillus subtilis* subsp. *subtilis* standard strain.