CASE REPORT

Meningoencephalitis Caused by a *Campylobacter fetus* in a Patient with Chronic Alcoholism

Sho Tanabe, Satoshi Kutsuna, Motoyuki Tsuboi, Nozomi Takeshita, Kayoko Hayakawa and Norio Ohmagari

Abstract:
We herein report a case of *Campylobacter fetus* meningoencephalitis in a patient with chronic alcoholism. *C. fetus* is a rare cause of meningitis. The patient presented with hallucinations and monology, and alcohol withdrawal was initially suspected. After he was unsuccessfully treated for alcohol withdrawal delirium, we diagnosed *C. fetus* meningoencephalitis. Ampicillin monotherapy gradually improved his clinical status. A previous report stated that *C. fetus* infection is associated with chronic alcoholism. In patients with chronic alcoholism and disturbed consciousness, an atypical bacterial central nervous system infection, such as *C. fetus* meningoencephalitis, should be considered.

Key words: *Campylobacter fetus*, meningoencephalitis, chronic alcoholism, ampicillin

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Introduction

Meningoencephalitis can cause disturbed consciousness, and rapid identification with prompt appropriate therapy is important for achieving optimal outcomes. However, this disease may be misdiagnosed because disturbed consciousness makes it difficult to obtain enough information from the patient. We herein report a case of *Campylobacter fetus* meningoencephalitis that was treated as alcohol withdrawal because of the patient’s history of habitual drinking.

Case Report

A 56-year-old man with a history of chronic alcoholism experienced disturbed consciousness over several days and was subsequently brought to the Psychiatry Department at our hospital from a nearby mental hospital. A psychiatric evaluation revealed that the patient presented with hallucinations and monology. Based on the patient’s history of chronic alcoholism, he was treated for alcohol-withdrawal-associated delirium. Upon admission, he also exhibited a fever and decubitus, which was treated with cefazolin (1 g every 8 hours) for presumed infection.

On day 6 of admission, positive results for *C. fetus* were obtained from his blood culture. Psychiatrists contacted our Department of Infectious Diseases and we initiated a careful evaluation of the patient. When we contacted the patient, he still had hallucination and monology. He did not complain of headache or nausea and did not have a history of eating raw meat before admission. A physical examination revealed normal findings of his respiratory system, cardiovascular system, and abdomen. Although he did not have a headache, we detected mild neck rigidity, possibly indicating meningitis. Laboratory tests also revealed a slightly elevated white blood cell count (9,910/μL) and C-reactive protein level (0.9 mg/dL).

A lumbar puncture was performed, and the patient’s cerebrospinal fluid (CSF) was clear and colorless. The initial pressure was 100 mmH₂O. The CSF cell count was 142/μL (neutrophils: 56/μL, lymphocytes: 86/μL), glucose level was 58 mg/dL (plasma glucose level 99 mg/dL, CSF to serum glucose ratio 0.57), and protein level was 94 mg/dL; no micro-organisms were observed on Gram staining of CSF.

Contrast-enhanced magnetic resonance imaging (MRI) revealed multiple areas of inflammation in the cortex, meninx, and dura mater, which indicated meningoencephalitis. Considering the results of the blood culture and contrast-
enhanced MRI, we started ampicillin (3 g every 6 hours) for suspected *C. fetus* meningoencephalitis. On day 16, we identified growth of *C. fetus* in the culture of a CSF specimen, and it took 10 days to receive the results of the culture after we obtained CSF. Our lab measured the minimum inhibitory concentration (MIC) by the E test for ampicillin (4 μg/ml). The patient’s condition gradually improved after initiating the ampicillin treatment, and we also detected improvements in his disturbed consciousness. After four weeks of intravenous ampicillin treatment, we observed that the patient had fully recovered his disturbed consciousness, and he was discharged. At three weeks post-discharge, MRI confirmed that his brain inflammation had improved (Figure).

**Discussion**

In the present case, the patient did not complain of typical meningitis symptoms, such as headache or nausea. Previous reports have shown that few *C. fetus* meningoencephalitis cases present with a classic meningitis triad (1). MRI was helpful in the diagnosis of *C. fetus* meningoencephalitis. The MRI finding implied central nervous system infection or limbic encephalitis and we could rule out other neurological diseases such as Wernicke’s encephalopathy.

Although *C. fetus* infections are most frequently observed among patients with immunocompromised status, such as those with diabetes, malignancy, corticosteroid therapy, or hepatic failure (2), we speculate that chronic alcoholism was strongly related to this case. In 1998, Dronda et al. reviewed the literature regarding *C. fetus* meningitis and reported that 4 of 8 cases (50%) involved chronic alcoholism (3). In addition, van Samkar et al. reviewed the literature regarding *C. fetus* meningitis and found that 9 of 22 patients (40%) with *C. fetus* meningitis between 1960 and 2013 had chronic alcoholism (1). In total, we identified 24 previously published cases that also showed a relationship between alcohol and *C. fetus* meningitis (Table). These findings imply that *C. fetus* meningoencephalitis may also be related to chronic alcoholism.

No treatment protocol for *C. fetus* meningoencephalitis has yet been established. While the present case was successfully treated with ampicillin monotherapy, as shown in Table, most previously documented *C. fetus* meningitis cases were treated with multiple or broad-spectrum antibiotics. However, there is no clear evidence that *C. fetus* meningitis should be treated with multiple antibiotics, as two other cases were successfully treated by ampicillin or amoxicillin monotherapy (4).

One other report recommended using imipenem for *C. fetus* central nervous system infections or patients with a severe status (5). However, this treatment has some problems. First, the use of broad-spectrum antibiotics such as

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![Figure.](image) **Figure.** Contrast-enhanced magnetic resonance image of the patient’s brain. (A) Fluid-attenuated inversion recovery images from before treatment revealed acute brain inflammation. (B) Images obtained at the three-week follow-up after discharge revealed improvement in the inflammation.
| Case no. (reference) | Sex | Age | Underlying Condition(s) | WBC in CSF | CSF Glucose | CSF Protein | Neck Stiffness | Headache | Confusion | Imaging | Treatment | Outcome |
|---------------------|-----|-----|--------------------------|------------|-------------|-------------|---------------|----------|-----------|---------|-----------|---------|
| Present case        | M   | 56  | Chronic alcoholism       | 142 μL     | 58 mg/dL    | 94 mg/dL    | +             | -        | +         | contrast MRI | ABPC     | Cured   |
| 1 (1) F             | 23  | NA  | NA                       | 308 μL     | 30.6 mg/dL  | 90 mg/dL    | +             | -        | -         | NA      | AMPC, CTRX | Cured   |
| 2 (1) M             | 52  | NA  | Chronic alcoholism       | 243 μL     | 18 mg/dL    | 176 mg/dL   | +             | -        | -         | NA      | AMPC, CTRX | Cured   |
| 3 (8) M             | 38  | NA  | Chronic alcoholism       | 2,048 μL   | 48.6 mg/dL  | 100 mg/dL   | +             | -        | -         | NA      | ABPC, GM   | Cured   |
| 4 (9) M             | 68  | NA  | Metastatic adenocarcinoma | NA         | NA          | NA          | NA            | NA       | NA        | NA      | ABPC, CTRX | Died    |
| 5 (9) M             | 65  | NA  | Alcoholic cirrhosis      | 48 μL      | 104.4 mg/dL | 90 mg/dL    | +             | -        | -         | NA      | CTRX, GM   | Cured   |
| 6 (10) M            | 47  | NA  | Diabetes mellitus        | 48 μL      | 81 mg/dL    | 90 mg/dL    | +             | -        | -         | NA      | CP, EM     | Cured   |
| 7 (11) F            | 39  | NA  | Chronic alcoholism       | 5.4 mg/dL  | 131 mg/dL   | 90 mg/dL    | +             | -        | -         | NA      | AMPC/CPVA | Cured   |
| 8 (11) M            | 36  | NA  | Chronic alcoholism       | NA         | 95 mg/dL    | 95 mg/dL    | +             | -        | -         | NA      | ABPC, THP  | Cured   |
| 9 (12) M            | 83  | NA  | Alcoholic cirrhosis      | 577 μL     | 91.8 mg/dL  | 60 mg/dL    | +             | -        | -         | NA      | CPFX, CTRX | Died    |
| 10 (3) M            | 47  | NA  | Alcoholic cirrhosis      | 300 μL     | 85 mg/dL    | 85 mg/dL    | +             | -        | -         | NA      | OFLX, GM   | Cured   |
| 11 (5) M            | 71  | NA  | Diabetes mellitus        | 11,100 μL  | 508 mg/dL   | 508 mg/dL   | +             | -        | -         | NA      | ABPC, GM, IPM, CTX, CPFX | Cured   |
| 12 (13) M           | 47  | NA  | Chronic alcoholism       | 2,128 μL   | 152 mg/dL   | 152 mg/dL   | +             | -        | -         | NA      | PEN, TET   | Cured   |
| 13 (14) M           | 55  | NA  | Chronic lymphocytic leukemia | 240 μL    | NA          | NA          | +             | -        | -         | NA      | PEN, TET   | Cured   |
| 14 (15) F           | 48  | NA  | Rheumatic fever          | 1,399 μL   | NA          | NA          | +             | -        | -         | NA      | PEN, CP    | Cured   |
| 15 (16) M           | 50  | NA  | Diabetes mellitus        | 3,436 μL   | 96 mg/dL    | 96 mg/dL    | +             | +        | -         | NA      | PEN, ABPC, CP | Cured   |
| 16 (17) F           | 69  | NA  | Diabetes mellitus        | 1,230 μL   | NA          | NA          | -             | -        | -         | NA      | PEN, CF, SFZ | Died    |
| 17 (18) M           | 40  | NA  | NA                       | 8,464 μL   | 120 mg/dL   | 120 mg/dL   | +             | -        | -         | NA      | ABPC, PEN, STR, EM | Died    |
| 18 (19) M           | 50  | NA  | NA                       | 36 μL      | 73 mg/dL    | 73 mg/dL    | +             | -        | +         | NA      | ABPC, CP    | Cured   |
| 19 (11) M           | 36  | NA  | Chronic alcoholism       | 156 μL     | 95 mg/dL    | 95 mg/dL    | -             | -        | -         | NA      | AMPC       | Cured   |
| 20 (20) M           | 55  | NA  | Alcoholic hepatitis      | 400 μL     | 83 mg/dL    | 83 mg/dL    | +             | -        | -         | NA      | ABPC       | Cured   |
| 21 (21) M           | 51  | NA  | NA                       | 344 μL     | 33 mg/dL    | 33 mg/dL    | +             | -        | -         | NA      | CTRX       | Cured   |
| 22 (22) M           | 40  | NA  | Crohn's disease          | 170 μL     | NA          | NA          | +             | -        | -         | NA      | CTRX, AZM  | Cured   |
| 23 (23) M           | 28  | NA  | Epilepsy                 | 170 μL     | NA          | NA          | +             | -        | -         | NA      | EM         | Cured   |
| 24 (4) M            | 75  | NA  | Diabetes mellitus        | 1430 μL    | 114 mg/dL   | 114 mg/dL   | -             | +        | -         | NA      | CTX, CPFX, AZM | Cured   |

ABPC: ampicillin, CTRX: ceftriaxone, GM: gentamicin, EM: erythromycin, CEZ: cefazolin, TOB: tobramycin, AMPC/CPVA: amoxicillin/clavulanate, CP: chloramphenicol, OFLX: ofloxacin, IPM: imipenem, CPFX: ciprofloxacin, PEN: penicillin, TET: tetracycline, SFZ: sulfadiazine, STR: streptomycin, MFLX: moxifloxacin, AZM: azithromycin
imipenem must be limited in order to prevent the growth of organisms resistant to antibiotics. Ohmagari et al. showed that the use of carbapenems for seven days was a risk factor for infections with multidrug-resistant *Pseudomonas aeruginosa* (6). Second, imipenem carries an additional risk. Van De Beek et al. recommended avoiding imipenem for meningitis treatment because it could lower the seizure threshold (7). In the present case, once positive results for *C. fetus* were obtained from the blood culture, we suspected *C. fetus* meningoencephalitis and started ABPC monotherapy. At this time, other bacteria were not suspected to be the cause of meningitis; therefore, we did not select empiric therapy and did not use imipenem, which is not recommended for bacterial meningitis.

When patients with chronic alcoholism present with central nervous system symptoms, we tend to diagnose alcohol-related diseases, such as alcohol withdrawal or Wernicke’s encephalitis. However, if the patient’s symptoms do not improve after appropriate treatment for alcohol withdrawal, we should consider the possibility of central nervous system infection and expand the examinations. As seen in the present case, results of MRI and lumbar puncture helped us make a correct diagnosis. In conclusion, we encountered a case of *C. fetus* meningoencephalitis with chronic alcoholism successfully treated by ampicillin monotherapy.

The authors state that they have no Conflict of Interest (COI).

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