Metronidazole-induced encephalopathy in a patient with Crohn’s disease

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Metronidazole is a widely used antibiotic for the treatment of anaerobic bacterial infections. Metronidazole-induced encephalopathy (MIEP) is a rare but potentially reversible disease. The mechanism of MIEP remains unclear, and differences in the neurotoxic effects of oral versus intravenous (IV) metronidazole administration have not yet been determined. We report the case of a Crohn’s disease (CD) patient who experienced encephalopathy immediately after a single IV dose of metronidazole following long-term exposure to the oral form of the drug. The 64-year-old man with intractable CD experienced a sudden change in mental status, aphasia, and muscle weakness after IV administration of metronidazole. He had previously taken metronidazole orally for 13 years and received intermittent IV metronidazole treatments for CD exacerbation. Brain magnetic resonance imaging (MRI) showed high-intensity signals in the bilateral medial thalamus and the midbrain and pontine tegmentum on fluid-attenuated inversion recovery images. After discontinuation of metronidazole, the high-intensity brain MRI signals resolved and the patient’s mental status dramatically improved; however, the patient exhibited mild cognitive dysfunction 2 months after the onset of encephalopathy. (Intest Res 2017;15:124-129)

Key Words: Metronidazole; Drug side effects; Metabolic encephalopathy; Crohn disease

INTRODUCTION

Crohn’s disease (CD) is an immunological disorder characterized by a relapsing-remitting clinical course. Antibiotics are effective in the treatment of CD, especially in patients with complicated fistulas or abscesses. Increasing evidence suggests that dysregulated gut microbiota play a role in the pathogenesis of CD. Metronidazole is a widely used antibiotic in patients with CD. However, metronidazole may elicit neurotoxic effects including peripheral neuropathy, ataxia, seizure, and encephalopathy. Metronidazole-induced encephalopathy (MIEP) is a very rare but potentially reversible disease. The exact mechanism of MIEP remains unclear. Here, we report the case of a CD patient who experienced encephalopathy immediately after a single intravenous (IV) dose of metronidazole following long-term exposure to the oral form of the drug.

CASE REPORT

A 64-year-old man who had been diagnosed with CD 17 years previously presented with altered mental status. He received sulfasalazine and azathioprine as maintenance treatment for CD and was intermittently treated with prednisolone to achieve remission after flare-ups. He had also received infliximab for the previous 3 years because of refractoriness to conventional treatment. Fourteen years before beginning infliximab treatment, the patient underwent ileocecectomy due to intractable active CD. Nine years after...
his first surgical treatment for active CD, he underwent right hemicolectomy and small bowel segmental resection to treat an intra-abdominal abscess communicating with multiple fistulas. He underwent three additional bowel resections because of a recurrent wound infection with enterocutaneous fistulas. The length of his remaining small intestine was 120 cm.

The patient had taken 250 mg of metronidazole orally three times daily beginning 13 years previously. After 8 years of treatment, however, he stopped taking metronidazole orally because of the side effect of peripheral neuropathy. He received IV metronidazole treatments three times over the following 5 years when he experienced complications such as enterocutaneous fistulas or abscesses. Six months before the last admission, however, he restarted and continued oral metronidazole treatment for active CD. The cumulative dose
of metronidazole over the 6 months prior to the last admission was 111.75 g. His lifetime cumulative dose of metronidazole was approximately 2,000 g.

The patient visited the hospital emergency department complaining of an abdominal wound problem which had been aggravated 2 days previously. He presented with pain, redness, and swelling surrounding the wound. Upon examination, he exhibited mild tenderness in the middle to lower abdomen. Initial laboratory findings showed elevated CRP (4.3 mg/dL), total bilirubin (4.9 mg/dL), and direct bilirubin (3.7 mg/dL), as well as prolonged PT (INR of 2.07). Other laboratory findings were as follows: white blood cell count 8,100/mm$^3$, hemoglobin level 12.4 g/dL, platelet count 140×10$^9$/L, cholesterol level 62 mg/dL, total protein level 5.6 g/dL, albumin level 2.8 g/dL, BUN level 10 mg/dL, serum creatinine level 0.57 mg/dL, serum AST level 22 IU/L, serum ALT level 16 IU/L, and serum ALP level 61 IU/L. Test results for all serological markers of hepatitis B and C were negative. Abdominal CT scans revealed a large enterocutaneous fistula in the middle portion of the patient’s abdomen (Fig. 1).

Considering the shortness of his small bowel, the patient received conservative treatment including parenteral nutrition and antibiotics for the treatment of the wound infection and enterocutaneous fistula. Metronidazole was administered intravenously during fasting.

Prior to IV metronidazole administration, the patient could follow commands and showed no motor weakness (Glasgow coma scale score of 15). However, he abruptly

![Fig. 3. Follow-up brain MRI findings. MR images obtained 7 days after metronidazole discontinuation. A fluid-attenuated inversion recovery image shows the improvement of lesions in the bilateral medial thalamus (A, B); midbrain tegmentum (C); pontine tegmentum, fourth ventricle, and dentate nucleus (D); and subcortical white matter (E).](image-url)
became comatose (Glasgow coma scale score of 8) 2 hours after receiving a single 500 mg IV dose of metronidazole. On neurological examination, the patient could not open his eyes upon painful stimulation but could localize the pain bilaterally. All brainstem reflexes were normal. However, bilateral Babinski signs were observed, suggestive of bilateral diffuse cortico-subcortical lesions. The pupillary light reflex was intact, and there was no limitation of extraocular muscle movement. The patient's vital signs were normal at the time of metronidazole administration, with a blood pressure of 139/82 mmHg, heart rate of 86 beats per minute, respiratory rate of 18 breaths per minute, and body temperature of 36.6°C. Two hours after IV metronidazole administration, however, his blood pressure was 160/96 mmHg and his heart rate was 133 beats per minute.

The rapid adrenocorticotropic hormone stimulation test revealed a normal response. The patient's brain CT scan was unremarkable. However, fluid-attenuated inversion recovery MRI showed multiple high-intensity lesions in the bilateral subcortical white matter, medial thalamus, midbrain and pontine tegmentum, bilateral middle cerebellar peduncles, and cerebellar dentate nucleus (Fig. 2). Based on clinical and radiological findings, a tentative diagnosis of MIEP was made. Metronidazole treatment was discontinued and replaced with IV piperacillin/tazobactam. A megadose of IV thiamine (500 mg) was administered three times daily for 8 days. The patient was subsequently maintained on 50 mg of thiamine daily.

Seven days after discontinuing metronidazole treatment, the patient was able to track moving objects. On day 8, a follow-up brain MRI showed more clearly visible diffuse bilateral subcortical white matter lesions. However, the extent of the previously noted hyperintense lesions was decreased in the thalamus, brainstem, and cerebellar dentate nucleus (Fig. 3). On day 12, the patient showed a significant improvement in his level of consciousness and was able to follow verbal commands and answer simple questions. Two months after the onset of MIEP, he was able to talk and walk unassisted. However, he still showed mild cognitive dysfunction. He received a score of 15 on the Korean version of the mini-mental state examination (normal range, 25–30) and demonstrated severe frontal dysfunction, as indicated by a score of 7 on the Frontal Assessment Battery (range, 0–18) (Fig. 4).

**DISCUSSION**

Metronidazole is associated with declines in CDAI scores and a reduced CD recurrence rate after surgery. It is well known that metronidazole is comparatively safe at the usual dosages. However, various neurological adverse events, including peripheral neuropathy, cerebellar dysfunction, and seizure, can develop. MIEP is one of the most serious adverse events associated with metronidazole treatment; however, it has a favorable prognosis after drug cessation. The long-term safety of large cumulative doses of metronidazole has not been elucidated. In previous reports, patients who experienced MIEP or cerebellar ataxia were not exposed to larger cumulative doses of metronidazole than that received by our patient; however, the association between neurotoxicity and the cumulative dose or duration of metronidazole treatment remains controversial.

Metronidazole-induced neurotoxicity typically occurs at doses exceeding 2 g/day for prolonged periods. Thus, our patient may have been at high risk of MIEP because of an extremely high cumulative dose of metronidazole received over a period of 13 years. Previous history of long-term exposure to metronidazole should be considered to prevent neurotoxic effects such as MIEP in patients who require repeated treatment.

Previous studies have shown that brain lesions identified by MRI in MIEP patients are typically located in the cerebellar dentate nucleus, dorsal medulla, dorsal pons, midbrain, and splenium of the corpus callosum, with bilateral and symmetrical involvement. Thus, the brain MRI findings in this case were consistent with changes associated with MIEP.

In this case, MIEP developed immediately after IV administration of a single dose of metronidazole following long-term oral metronidazole treatment. Differences in the neurotoxic effects of oral versus IV metronidazole adminis-
traction are not yet understood. A previous study showed that the upper part of the small intestine is the main site of metronidazole absorption.\textsuperscript{12} Manafi et al.\textsuperscript{13} found that absorption of orally administered metronidazole was significantly reduced by postoperative ileus. Therefore, a short bowel may be related to poor absorption of both nutrients and the metronidazole. The peak serum level of metronidazole after IV administration in patients with a short bowel may be higher than that after oral administration.

Because metronidazole is mainly metabolized in the liver, severe liver dysfunction increases the drug’s half-life and increases the risk of MIEP.\textsuperscript{14} In our case, there were no clinical signs suggesting chronic liver disease or hepatic encephalopathy, although the patient had cholestasis and PT prolongation at admission. In addition, the abdominal CT scan showed neither cirrhotic changes nor mechanical biliary obstruction. Thus, these transient abnormalities revealed by the liver function test may have been related to impaired hepatocyte function and systemic infection caused by bowel inflammation and bacterial overgrowth.\textsuperscript{15} Liver function test results were stable for 2 weeks after admission and gradually returned to normal. In contrast, the patient's mental status began to improve within a week after cessation of metronidazole treatment. This temporal relationship supports the diagnosis of MIEP rather than hepatic encephalopathy.

In this case, Wernicke's encephalopathy (WE) may be another possible diagnosis, as acute WE has clinical symptoms and brain MRI findings similar to those of MIEP. Metronidazole metabolites inhibit thiamine pyrophosphokinase, and the resulting thiamine antagonism may also play a role in the pathogenesis of MIEP.\textsuperscript{16} However, the patient displayed altered mental status immediately after IV administration of metronidazole. The temporal relationship between metronidazole administration and onset of neurological symptoms suggests a diagnosis of MIEP rather than WE. Moreover, the patient showed no clinical signs of oculomotor dysfunction such as nystagmus and conjugate gaze palsy, which are typical clinical features of WE.\textsuperscript{17} Brain MRI revealed diffuse high-intensity lesions in the bilateral subcortical white matter, medial thalamus, midbrain and pontine tegmentum, bilateral middle cerebellar peduncles, and cerebellar dentate nucleus, which are compatible with a diagnosis of MIEP. Lesions in the dentate nucleus and red nucleus can also be found in patients with WE, although these findings are atypical.\textsuperscript{18} The most common structures affected by WE, as demonstrated by MRI, are the medial thalamus and periventricular regions of the third ventricle.\textsuperscript{19,20} In a systematic review of 64 patients with MIEP, MRI revealed subcortical white matter lesions in approximately 50% of patients with altered mental status.\textsuperscript{9} However, high-intensity lesions in the bilateral subcortical white matter have not been detected in patients with WE.\textsuperscript{9,11,16-20} These findings strongly suggest that MIEP is the primary cause of altered mental status in this case. Nonetheless, the possibility of concomitant WE could not be completely excluded because the patient was at high risk of thiamine deficiency due to malnutrition resulting from his short bowel status and was treated with thiamine. Moreover, the subtle cardiovascular signs observed in our patient, such as tachycardia, are frequently also observed in patients with acute WE. Because the patient's serum thiamine level was not measured before thiamine replacement therapy, the possibility of concomitant WE may be a limitation in this case study.

We report the first case of a patient with CD who experienced encephalopathy immediately after IV administration of a single dose of metronidazole following long-term oral metronidazole treatment. MIEP may be a rare complication in CD patients, especially those at risk of malnutrition or who have received a high cumulative dose of metronidazole over a long period. Therefore, evaluation of nutritional status is very important in patients with IBD to prevent irreversible complications such as WE.\textsuperscript{11,22}

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