Intrathecal Isoniazid for Refractory Tuberculous Meningitis with Cerebral Infarction

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Abstract

A 30-year-old Vietnamese woman, about 19 weeks pregnant, was admitted for acute cerebral infarction with stenosis of the left middle cerebral artery (LMCA), tuberculous meningitis, and miliary tuberculosis. Treatment with heparin, quadruple anti-tuberculosis therapy, and dexamethasone afforded prompt symptomatic improvement. However, she delivered a stillbirth, after which there was recurrence of acute cerebral infarction with LMCA occlusion, sinus thrombosis, and cranial base inflammation. A thrice-weekly 100 mg dose of intrathecal isoniazid (INH) improved the signs of meningeal inflammation. The patient was discharged ambulatory after 7 months. In refractory tuberculous meningitis, multimodal therapy with intrathecal INH and steroids should be considered.

Key words: intrathecal isoniazid, tuberculous meningitis, cerebral infarction, pregnancy

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Introduction

Tuberculous meningitis is a serious disease with high rates of mortality and residual neurologic deficits (1). Although the British Infection Society has recommended treatment regimens (2), additional high-strength modalities may be needed in some serious cases. We herein report a case of tuberculous meningitis in which remission was achieved not with the recommended treatment but with intrathecal administration of isoniazid (INH).

Case Report

The patient was a 30-year-old Vietnamese woman with no relevant medical history who was studying in Japan. She presented with common cold symptoms for 2 weeks and a fever of about 38°C, followed by right hemiplegia. These prompted consultation and subsequent admission to our hospital. The patient had undergone fertility treatment at a private obstetrics-gynecology clinic and was about 19 weeks pregnant. She had hyperemesis with a poor nutritional status, however, no weight gain during pregnancy was observed.

The general physical findings were as follows: height 157 cm, weight 52 kg, body mass index 21.1, blood pressure 116/57 mmHg, heart rate 128/min, respiration rate 30/min, body temperature 38.0°C, and oxygen saturation of 95%. There were no pertinent thoracic or abdominal findings, except for the signs of pregnancy. The neurologic findings were lucid consciousness, spontaneous pain in the posterior cervical region, nuchal rigidity, right central facial palsy, right hemiplegia (Brunnstrom stage I in the arm and fingers, Brunnstrom stage II in the leg), hyperreflexia of the right upper and lower limbs, and right leg sensory deficit. Blood inflammatory markers showed a white blood cell count of 6,920/μL and C-reactive protein of 3.42 mg/dL. Blood biochemistry results indicated mild liver dysfunction (aspartate aminotransferase 43 IU/L, alanine aminotransferase 46 IU/L) and undernutrition, with albumin at 2.3 g/dL. There were no abnormal findings related to the kidney or pancreas function. Testing for human immunodeficiency virus was negative.

The cerebrospinal fluid (CSF) findings were as follows:
pale yellow color, initial pressure 290 mmH₂O, final pressure 150 mmH₂O, cell count 100/μL with a lymphocyte-to-polymorphonuclear leukocyte ratio of 67:33, protein 229 mg/dL, sugar 11 mg/dL, and positive Ziehl-Neelsen staining. Cranial magnetic resonance imaging (MRI) on diffusion-weighted images revealed acute cerebral infarction of the left putamen; magnetic resonance angiography (MRA) revealed stenosis of the left middle cerebral artery (LMCA) (Fig. 1a and b). Ground glass opacities in both lung fields were noted on chest radiograph and computed tomography (CT) (Fig. 1c and d). Cerebral infarction secondary to vasculitis from tuberculous meningitis and miliary tuberculosis were suspected.

Fig. 2 shows the clinical course of the patient’s hospitalization. Treatment was started with dexamethasone (19.6 mg/day) and four anti-tuberculosis drugs: INH (200 mg/day), rifampicin (RFP) (450 mg/day), ethambutol (EB) (750 mg/day), and pyrazinamide (PZA) (1.2 g/day). Because bacterial meningitis could not be ruled out, meropenem (6 g/day) was also administered. Taking into consideration the patient’s pregnancy, acute cerebral infarction was treated with continuous heparin infusion at 10,000 units/day. Based on positive tubercle bacilli DNA on polymerase chain reaction and positive culture for Mycobacterium tuberculosis, the patient was given a definitive diagnosis of tuberculous meningitis. After five days of treatment, liver dysfunction ensued. When the liver dysfunction improved after a week, the administration of the four drugs was continued, with PZA being replaced by levofloxacin (LVFX) (500 mg/day). The fever and right hemiplegia promptly improved. Although Ziehl-Neelsen staining was performed only once and showed positivity and susceptibility to all anti-tuberculous drugs, the presence of Mycobacterium tuberculosis in the CSF was sustained for up to one month, despite the administration of anti-tuberculosis drugs.

On hospital day 25, at 22 weeks and 1 day of gestation, the patient gave birth to a stillborn. Examination of the uterine contents revealed tubercle bacilli, confirming an intrauterine infection. After the stillbirth, the fever and meningitis signs recurred. Despite treatment with 5 anti-tuberculosis agents (with the addition of streptomycin at 1 g/day) and restarting dexamethasone administration (19.6 mg/day), no improvement in the signs of meningeal irritation was observed. On hospital day 33, aphasia and right hemiplegia appeared; on hospital day 37, right abducens nerve palsy developed. Repeat cranial MRI revealed recurrent cerebral infarction on the left parietal lobe; MRA showed peripheral occlusion of the horizontal segment of the LMCA. Findings indicative of inflammation of the cranial base and lateral sinus thrombosis were also observed (Fig. 3). These were treated with
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Figure 2. Clinical course of a 30-year-old woman with refractory tuberculous meningitis. INH: isoniazid, RFP: rifampicin, EB: ethambutol, PZA: pyrazinamide, LVFX: levofloxacin, SM: streptomycin

tuberculosis drugs

Figure 3. Cranial MRI findings of recurrent cerebral infarction in tuberculous meningitis. (a) Diffusion-weighted images show infarcted area in the left posterior limb of the internal capsule, left corona radiata, and left parietal lobe. (b) MRA shows occlusion of the left middle cerebral artery (arrow head). (c) Enhanced T1-weighted images show inflammatory changes in the base of the brain (encircled area) and venous sinus thrombosis (arrow). MRI: magnetic resonance imaging, MRA: magnetic resonance angiography
edaravone and heparin; administration of aspirin (100 mg/day) was also continued to prevent recurrence.

However, the meningoits symptoms persisted and did not improve despite increasing the INH dosage from 300 mg/day to 500 mg/day. Therefore, intrathecal administration of 100 mg of INH 3 times per week was started. Furthermore, 15 mL of CSF was intermittently drained by lumbar puncture to relieve intracranial pressure. The signs of meningeal irritation started to improve after starting intrathecal INH. Serial cranial MRI indicated resolution of the venous sinus thrombosis and reduction of the inflammatory findings in the cranial base (Fig. 4). On hospital day 98, the visual acuity on the left decreased, suggesting left optic neuritis caused by inflammation of the cranial base. As the possibility of optic neuropathy could not be ruled out, EB was discontinued. Intrathecal INH administration was tapered to once a week and eventually was discontinued as the patient’s symptoms improved.

The patient was discharged for home on hospital day 191. Three anti-tuberculosis drugs (INH, RFP, and LVFX) were continuously administered for 1 year. The residual neurologic deficits were mild aphasia, decreased visual acuity on the left side, and hypoesthesia in both legs, which was suspected to be secondary to INH. Nevertheless, the patient was able to independently engage in her activities of daily living. Cranial MRI 2 years after onset showed that the occlusion in the horizontal segment of the LMCA was unchanged; therefore, the administration of aspirin (100 mg/day) was continued to prevent the recurrence of cerebral infarction. Approximately 2.5 years later, the patient returned to her home country. She was able to walk without assistance and carry out her usual duties and activities.

**Discussion**

Tuberculous meningitis causes inflammatory changes in the base of the brain, leading to vasculitis, cranial polyneuritis, and obstructive hydrocephalus (3). Cerebral infarction as a complication is widely known to be associated with a poor prognosis (4).

In the present case, we referred to the recommendations of the British Infection Society (2) and administered a four-drug combination of INH, RFP, EB, and PZA. Although the guidelines for treating tuberculosis in pregnant patients state that PZA should be avoided as much as possible because its safety has not been established (5), we decided to continue PZA to save the mother’s life, as this was a severe case of tuberculosis. Based on a report by Thwaites et al. (6), we also administered corticosteroids. The recommended treatment led to temporary improvement in the symptoms. However, this was a severe case of recurrent meningitis and cerebral infarction, sinus thrombosis, and cranial neuropathy, for which sustained improvement was only achieved by additional intrathecal administration of INH.

There have been only a small number of reports on the use of intrathecal INH. Marked improvement was noted in a case of refractory tuberculous meningitis after thrice-weekly intrathecal administration of 100 mg/day of INH (7). This therapy was also proven to achieve remission in a patient with tuberculous meningitis and liver dysfunction, where the oral administration of high-dose INH was not possible (8). In other reports, two patients were able to return to society without any serious residual effects after combination of intrathecal INH and steroid pulse therapy (9). These findings suggest that intrathecal administration of INH may contribute to an improved prognosis in refractory tuberculous meningitis.

After administration, INH rapidly diffuses into the CSF and reaches peak concentrations in excess of 3 mg/L, which is over 30 times its minimal inhibitory concentration (MIC) against *Mycobacterium tuberculosis*, within 4 hours (10, 11). INH is metabolized in the liver via the N-acetyltransferase 2 (NAT2), which is a polymorphic gene with enzymatic activity that can be altered by the presence of single nucleotide
polymorphisms (SNPs) in its coding region (12). Three major genetically determined phenotypes are observed: rapid, intermediate, and slow acetylators (12, 13). Among Brazilian patients with tuberculosis, 72.8% were found to be fast acetylators (14). A meta-analysis showed a significant association between the slow acetylator genotype of NAT2 and antituberculosis drug-induced hepatotoxicity; significant results were also found in East Asians, South Asians, Brazilians, and Middle Eastern populations, but none in Caucasians (15). This implies that the efficacy of INH may be based on NAT2 gene polymorphism and race.

In our case, the resistance to standard tuberculous therapy and the effectiveness of intrathecal INH were probably due to 1) disturbance of CSF circulation, 2) decline in INH penetration to the focal lesion, and 3) the possibility that the patient was a rapid acetylator of NAT2, although we did not investigate the polymorphism status in the present case. Overall, these findings suggest that refractory tuberculous meningitis or tuberculous meningitis complicated by anti-tuberculosis drug-induced hepatotoxicity may be indications for treatment with intrathecal INH.

This case exhibited recurrent meningitis symptoms and required administration of corticosteroids over several months. Corticosteroids can improve CSF findings, reduce intracranial pressure, and decrease inflammation (16, 17). It has also been suggested that corticosteroids be routinely used in HIV-negative people with tuberculous meningitis to reduce the risk of death and disabling residual neurologic deficit among survivors (1). Thwaites et al. (6) reported a protocol involving steroid discontinuation after 4 weeks. However, in refractory cases with persistent meningitis symptoms longer than 4 weeks, as exemplified by our patient, we believe that prolonged administration of corticosteroids can be considered.

In conclusion, multimodal therapy, including intrathecal administration of INH and systemic steroids, should be considered for cases of refractory tuberculous meningitis.

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

References


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