A 37-year-old woman presented to a community hospital with a 1-week history of severe frontal headache, subjective fever and episodic confusion. She denied having nausea, vomiting, seizures or rash. She had immigrated to Canada from Nigeria 8 months before presentation, with no travel since, and had no sick contacts or animal exposures.

On admission, the patient was febrile (38.3°C), but vital signs were otherwise unremarkable. She had nuchal rigidity without focal neurologic deficits. Bloodwork showed a normal leukocyte count and an absolute lymphocyte count of 0.7 (normal 1.0–3.57) × 10^9 cells/L. Test results for HIV were negative, as were blood smears for malaria parasites. Initial cerebrospinal fluid (CSF) studies showed a leukocyte count of 96 (normal 0–5) × 10^6 cells/L (59% lymphocytes, 36% neutrophils and 2% monocytes), glucose of 1.9 (normal 2.2–3.9) mmol/L and protein of 1.20 (normal < 0.45) g/L. At admission, her aspartate aminotransferase was 13 (normal 10–40) U/L, and her alanine aminotransferase was 18 (normal 5–42) U/L. Results from CSF bacterial and fungal cultures, polymerase chain reaction for herpes simplex virus and enterovirus, cryptococcal antigen lateral flow assay and autoimmune encephalitis work-up were negative. Mycobacterial culture of the CSF was not requested.

Concern for meningoencephalitis prompted a noncontrast computed tomography (CT) scan of the patient’s head, which showed a small area of subcortical hypodensity in the right temporal lobe, suspicious for encephalitis. Her clinicians started empiric treatment with ceftriaxone, vancomycin and acyclovir. On day 5, she became nonverbal and confused, and developed a new left lateral rectus palsy and an ipsilateral, nonreactive pupil measuring 6 mm. Brain magnetic resonance imaging showed high signal intensity throughout the convexity sulci and basal cisterns, compatible with meningitis (Figure 1). A CT scan of the chest, abdomen and pelvis showed nonspecific abdominal lymphadenopathy. She developed Cushing reflex and was intubated for transfer to our tertiary hospital for neurosurgical intervention. Neurosurgeons inserted an external ventricular drain to relieve hydrocephalus, and escalated antimicrobial therapy to meropenem, vancomycin and acyclovir.

Figure 1: Magnetic resonance imaging (MRI) scan of the brain of a 37-year-old woman with tuberculous meningitis, obtained on day 5 of hospital admission. This axial fluid-attenuated inversion recovery (FLAIR) image at the level of the basal cisterns and temporal lobes shows high signal intensity throughout the convexity sulci and basal cisterns, compatible with meningitis. Both temporal horns of the lateral ventricles are dilated, with abnormal signal intensity surrounding them, compatible with hydrocephalus and transependymal passage of cerebrospinal fluid. The initial MRI report commented on possible parenchymal abnormality in the right temporal lobe (yellow arrow), which was thought to be compatible with encephalitis. Upon review of this image, however, the MRI findings are most compatible with meningitis.

Key points
• In Canada, tuberculous meningitis is an uncommon presentation of Mycobacterium tuberculosis infection, occurring most frequently in individuals who arrived to Canada within the previous 5 years.
• In patients with risk factors for tuberculosis (TB), a presumptive diagnosis of TB meningitis should be considered if clinical presentation and findings from cerebrospinal fluid are consistent; early empiric therapy should be initiated.
• Clinicians should monitor patients closely for adverse reactions to antituberculous medications, such as transaminitis and acute kidney injury, and stop potentially hepatotoxic drugs, where possible.
• Patients with chronic hydrocephalus as a complication of their meningitis may require a protracted course of corticosteroids, in addition to extraventricular drainage.

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Meningitis caused by Mycobacterium tuberculosis in a recent immigrant to Canada

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We consulted the infectious diseases team, who considered Mycobacterium tuberculosis meningitis to be the primary differential diagnosis given the patient’s risk factors, progressive presentation despite treatment with antimicrobial agents, CSF biochemical findings and abdominal lymphadenopathy. We stopped antimicrobial therapy, performed another lumbar puncture to obtain CSF for mycobacterial culture and GeneXpert MTB/RIF nucleic acid amplification testing (NAAT; Cepheid), and started empirical treatment for tuberculous (TB) meningitis with intravenous (IV) dexamethasone 0.4 mg/kg/d divided every 4 hours, IV rifampin 10 mg/kg/d, oral isoniazid 5 mg/kg/d, oral pyrazinamide 25 mg/kg/d and oral ethambutol 15 mg/kg/d. Acid-fast bacilli were not seen on CSF staining, and NAAT was negative for M. tuberculosis complex. Our intensivists performed bronchoscopy, from which NAAT, staining and culture were ultimately negative.

Beginning on day 18, adverse events necessitated a series of changes to the patient’s treatment regimen (Figure 2). Her levels of aspartate aminotransferase (141 U/L) and alanine aminotransferase (227 U/L) were more than 5 times the upper limit of normal, which prompted us to stop pyrazinamide and isoniazid. We substituted with IV amikacin 15 mg/kg/d and IV levofloxacin 10 mg/kg/d. Amikacin caused acute kidney injury, with a doubling of serum creatinine over 4 days, so we switched to oral linezolid 600 mg twice daily. Once transaminase levels returned to normal, we stopped linezolid and reintroduced isoniazid. As the patient later developed a transaminitis that we attributed to isoniazid, we permanently stopped isoniazid and reintroduced linezolid.

M. tuberculosis was detected in liquid culture of CSF on day 37. The isolate was negative for mutations causing resistance to rifampin and isoniazid, and the laboratory later confirmed susceptibility to first-line (i.e., isoniazid, rifampin, pyrazinamide, ethambutol) and second-line (i.e., ethionamide, linezolid, moxifloxacin) agents.

Our initial attempts to taper the patient off of dexamethasone according to routine schedules were unsuccessful because of persistent hydrocephalus; she continued dexamethasone treatment during her hospitalizations and was switched to hydrocortisone upon discharge to allow for a slow taper of steroids over 5 months (Figure 3). The patient’s external ventricular drain was weaned and eventually removed on day 31. On day 35, she was extubated; shortly thereafter, however, neurosurgeons reinserted an external ventricular drain for worsening hydrocephalus and ultimately exchanged it for a programmable ventriculoperitoneal shunt (VPS). This was later replaced with a medium-pressure valve VPS after a short period of worsening hydrocephalus and generalized myoclonus concerning for seizures. The patient was eventually discharged home on day 148 from the community hospital.

The patient remained stable at 1-year follow-up, and her antituberculous medications were stopped. Though she recovered functional independence and regained verbal communication skills, she continues to suffer from disease sequelae and treatment-related complications, particularly cerebral salt wasting syndrome and bilateral optic nerve damage, and has been unable to return to work.

Discussion

Tuberculosis remains a leading cause of death globally, with 1.4 million deaths in 2019. In Canada, most reported cases of TB are among foreign-born individuals, with more than 40% occurring within the first 5 years of arrival. Tuberculosis of the central ner-
you system accounts for 1%–2% of all TB cases, causing a spectrum of disease that includes brain abscess, cerebritis, meningitis, myelitis and tuberculomas. Tuberculous meningitis is the most common central nervous system manifestation, with substantial morbidity (25%) and mortality (15%–40%) rates, even with appropriate treatment. Independent risk factors for death among inpatients with TB meningitis include HIV infection, age older than 40 years, CSF culture positive for TB and disease severity. Tuberculous meningitis typically presents subacutely, with fever, headache, malaise and personality changes, which can progress to cranial nerve palsies or other focal neurologic deficits, seizures, coma and death. Lumbar puncture is vital, as CSF findings can be suggestive of a diagnosis of TB meningitis and prompt the decision to start empiric therapy. Typical CSF findings include an elevated opening pressure, marked lymphocytic pleocytosis, elevated protein caused by obstructed CSF flow, and low glucose. Microbiologic testing is confirmatory in the diagnosis of TB meningitis, but it has limitations. Samples of CSF should be sent for acid-fast bacillus staining and microscopy, NAAT and culture. However, the mycobacterial burden in CSF is generally lower in patients with TB meningitis than with pulmonary TB, which affects the yield from microbiologic testing. The sensitivity of staining for acid-fast bacillus and microscopy (20%–80%) is lower in patients with extrapulmonary TB and in patients with HIV infection; thus, a negative result from acid-fast bacillus staining of CSF should not lead clinicians to rule out TB meningitis. However, the sensitivity of staining for acid-fast bacillus in TB meningitis can be increased by performing daily lumbar punctures for 3 days, submitting the last tube collected and collecting a large sample volume (10–15 mL). Similarly, NAAT testing, which has the advantage of providing a presumptive diagnosis if positive, with a turnaround time of less than 24 hours, has a lower sensitivity in CSF than sputum. Concomitant pulmonary TB is found in 22%–24% of patients with TB meningitis, and sputum examination offers a less invasive diagnostic modality; therefore, patients with presumed TB meningitis should be investigated for pulmonary TB. In our case, work-up for pulmonary TB was negative. Mycobacterial cultures can take up to 7 weeks and a negative culture cannot exclude TB meningitis in patients without an alternative diagnosis who have improved on antituberculous therapy. Therefore, if clinical presentation and CSF findings are consistent, a presumptive diagnosis of TB meningitis should be considered for patients with risk factors for TB. It is imperative that early empiric therapy is initiated, as time to initiation of appropriate antituberculous therapy before coma onset is the strongest predictor of survival.

Current guidelines recommend 2 months of intensive phase treatment with rifampin, isoniazid, pyrazinamide and ethambutol for drug-susceptible TB, followed by continuation phase treatment with rifampin and isoniazid for an additional 10 months. No difference in clinical outcome or relapse has been found with a 6-month treatment duration. Rifampin and ethambutol remain cornerstones of intensive phase therapy for TB meningitis because small trials have failed to show an improvement in mortality rates when better CSF-penetrating agents are added.
As shown with our patient, antituberculous therapy requires careful monitoring for adverse events. When encountered, identification of the most likely causative agent, a decision about its temporary cessation or permanent discontinuation, and choice of an acceptable alternative is important. Clinicians should consider second-line agents with acceptable CSF pharmacokinetic parameters, such as levofloxacin, linezolid, moxifloxacin, cycloserine or ethionamide. These parameters led us to choose a twice daily linezolid dosage during the patient’s critical illness while susceptibility testing results were unavailable, as opposed to the once daily dosage typically used in trials of TB meningitis. One study showed improvement in CSF biochemical parameters and Glasgow Coma Scale with the regimen we used.

Potentially hepatotoxic drugs should be stopped when serious transaminitis occurs. Guidelines recommend stopping pyrazinamide, isoniazid and rifampin, and substituting with either a fluoroquinolone and an injectable agent (e.g., aminoglycosides, capreomycin), or a fluoroquinolone and an oral second-line agent (e.g., linezolid). We withheld the 2 most likely causes of transaminitis, isoniazid and pyrazinamide, but continued rifampin to avoid stopping most first-line agents during our patient’s critical illness.

Successful treatment of TB meningitis requires a multimodal approach to manage increased intracranial pressure, including corticosteroids, which have been shown to reduce risk of death. The optimal dose and duration of corticosteroids is unclear, with guidelines suggesting a 6-8 week taper. Our case highlights the limitation of these recommendations and the importance of tailoring an individual’s therapy, as multiple attempts to wean the patient’s corticosteroids failed; she ultimately required a 48-week taper.

Management of associated hydrocephalus, especially if noncommunicating, may require urgent neurosurgical intervention such as an external ventricular drain for short-term control and — as in our patient — a VPS for definitive management. Optic nerve complications are responsible for most vision loss associated with TB meningitis and can have multiple causes, but were attributed to chronic hydrocephalus in our patient given the time of onset. Hyponatremia can be caused by cerebral salt wasting syndrome or syndrome of inappropriate antidiuretic hormone secretion, which need to be distinguished because of their different treatment approaches. Our patient’s hyponatremia was attributed to cerebral salt wasting syndrome, and corrected with oral supplementation of sodium chloride.

Our case shows the diagnostic and therapeutic challenges of managing TB meningitis. Early recognition and institution of appropriate therapy, with continued monitoring throughout the disease course, maximizes the probability of functional recovery.

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The section Cases presents brief case reports that convey clear, practical lessons. Preference is given to common presentations of important rare conditions, and important unusual presentations of common problems. Articles start with a case presentation (500 words maximum), and a discussion of the underlying condition follows (1000 words maximum). Visual elements (e.g., tables of the differential diagnosis, clinical features or diagnostic approach) are encouraged. Consent from patients for publication of their story is necessary. See information for authors at www.cmaj.ca.

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