Nausea, vomiting and diarrhea: An unusual presentation of multiple sclerosis

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Multiple sclerosis is a multifocal demyelinating disorder occurring with highest incidence in temperate climate zones. It usually affects young adults with subacute onset of focal neurological symptoms. The majority (80%) of patients present with visual, sensory or gait disturbances (1,2), whereas older patients (older than 40 years) more commonly exhibit symptoms of progressive myelopathy (2).

Nausea, vomiting and diarrhea are three of the most common symptoms encountered in patients seen by both family physicians and gastroenterologists. In the vast majority, the causes include specific or nonspecific gastrointestinal infections (3). However this symptom complex can be caused by neurological disease. The usual neurological etiologies include autonomic neuropathies (4,5), but rarely central causes can be seen (6-8). This symptom complex is an unusual presenting feature and may easily confound the diagnosis of multiple sclerosis, particularly if focal central nervous system signs are absent. The case of a young woman whose initial symptoms of nausea, vomiting and diarrhea appear to represent a first attack of multiple sclerosis is presented.

CASE PRESENTATION

In June 1990 a 33-year old right-handed woman presented to the emergency room with a two-week history of nausea, vomiting and intermittent (one to four times/day) loose watery stools without blood. The vomiting and diarrhea were not associated with cramps, fever or chills. She experienced mild light-headedness, especially on arising, and complained of mild left neck pain. On the day of admission she vomited three times. There were no symptoms of upper respiratory tract infection and no history of travel or contact with subjects with diarrhea. She is married and had two healthy children by caesarian section. She was last treated with antibiotics for sinusitis in 1985. There was no history of alcohol or substance abuse. The patient’s mother died of disabling multiple sclerosis, and a younger brother was recently diagnosed with this demyelinating disease.

Physical examination at admission disclosed a thin woman in no acute distress, with normal vital signs and no postural changes. The general and neurological examinations were entirely within normal limits.
Hematocrit was 36.3%, white blood cell count $5.3 \times 10^9/L$, mean cell volume 85.6 fl, blood glucose 4.5 mmol/L, blood urea nitrogen 2.9 mmol/L and creatinine 84 $\mu$mol/L. Electrolytes and liver enzymes were normal. A free thyroxine index and thyroid stimulating hormone were normal. Urinalysis and urine cultures were normal. A Venereal Disease Research Laboratory test for syphilis was negative. An electrocardiogram test was normal.

The patient was admitted for further evaluation. A gastroenterological work-up consisting of stool occult blood, cultures, sensitivity, ova and parasites were all negative. Gastroscopy and an upper gastrointestinal small bowel follow-through study were also normal. A colonoscopy to splenic flexure with biopsy of normal-looking mucosa disclosed mild nonspecific edema and occasional inflammatory cells in the rectosigmoid. A liquid gastric emptying scan showed 73% retention after 30 mins (normal less than 50%). A solid gastric emptying scan was normal.

Because symptoms persisted and nausea was a prominent feature, and because of the family history, neurological consultation was obtained. Clinical evaluation and a Bárány test were normal. A noncontrast computed tomographic (CT) scan of the head revealed a small hypodense lesion in the left subcortical white matter. A follow-up double dose delayed contrast CT scan of the head revealed two diffusely enhancing left parietal white matter lesions compatible with multiple sclerosis. Magnetic resonance imaging (MRI) with $T_2$-weighted images disclosed a much wider involvement of the subcortical white matter, with lesions in the left parietal, left occipital, right parietal and right optic radiation, as well as a lesion in the corpus callosum (Figure 1). A lumbar puncture revealed normal cells, protein and glucose, and negative bacterial microbiology. However, oligoclonal bands were present, supporting the diagnosis of multiple sclerosis. Somatosensory evoked potentials showed interference with conduction in the left cerebral hemisphere consistent with lesions in the left subcortical white matter.

The patient improved somewhat on dimenhydrinate suppositories and was discharged after eight days of hospitalization. However, she was readmitted with worsening nausea and vomiting six days later; she also complained of diffuse nonthrobbing headaches. She was started on intravenous methylprednisolone sodium 250 mg q6h and intravenous promethazine hydrochloride 25 mg q4h. During this second hospitalization, an episode of right hand and right thigh paresthesia lasted two to three days. A repeat CT scan of the head showed a prominent demyelinating plaque in the left corticomedullary region. She gradually improved and was discharged on oral promethazine hydrochloride 26 days after the second hospitalization. Promethazine hydrochloride was discontinued three months later, without a relapse of gastrointestinal symptoms. She has been well without gastrointestinal or neurological symptoms for the past five years.

**DISCUSSION**

Gastrointestinal manifestations of multiple sclerosis include transfer and transport dysphagia (9), nausea, vomiting (2), Brûn's syndrome (vertigo, vomiting, headache and visual disturbances with positional change of the head) (2,10), constipation and fecal incontinence (11,12). Inflammatory
bowel disease has also been associated with multiple sclerosis (13). In the majority of reports gastrointestinal symptoms are described in patients with established multiple sclerosis and most often are accompanied by objective neurological findings. Gastrointestinal work-up is usually negative.

As a primary initial manifestation of multiple sclerosis, nausea and vomiting are distinctly unusual and are most commonly associated with vertigo and eighth nerve involvement. Diarrhea, except in the context of fecal incontinence, is not mentioned as a feature of multiple sclerosis.

The initial diagnostic impression in our patient was that of prolonged gastroenteritis. However, the prominence of nausea was felt to be disproportional to other features. The subsequent gastroenterological work-up did not disclose any plausible explanation for the patient’s symptoms. Her family history, amalgamated neurological evaluations, MRI findings (14) and cerebrospinal fluid abnormalities (oligoclonal banding) established that our patient had familial multiple sclerosis. The explanation of her symptoms, however, is less clear. Overt demyelinating plaques were not visualized in the usual centres associated with vomiting. Lesions involving the dorsal motor nucleus of the vagus, the nucleus ambiguous or the autonomic nuclei in the medullary reticular formation are classically associated with vomiting and upper gut motility disturbances (15,16). Lesions in the floor of the fourth ventricle, the area postrema or chemotactic trigger zone for vomiting can lead to similar symptoms (8). It is, however, not an absolute requirement for diagnosis that one of the specific centres described be involved directly. Afferent or efferent fibre connections to these regions may mediate similar symptoms. For example, the lateral tegmentum of the pons and middle cerebellar peduncle have been implicated in the projectile vomiting of a man with metastatic malignant melanoma (16). However, microscopic plaques below the resolution of MRI or CT scanners in relevant areas cannot be ruled out and may have played a significant role in her symptoms.

In contrast to the dysautonomia commonly observed in diabetes mellitus (18,19,20), the gastric emptying disorder in patients with central causes may respond poorly to prokinetic therapy. The gastric emptying disturbance, both in the present case and in a previously reported patient with prolonged nausea, vomiting and diarrhea secondary to a posterior fossa tumour (7), did not respond to such therapy.

The diarrhea experienced by our patient is more difficult to explain. The presence of edema and spotty inflammatory cells on sigmoid biopsy raised the possibility of an unrecognized infectious agent. However, no agent was found and the upper gut symptoms were much more prominent than diarrhea. We speculate that her diarrhea and upper gastrointestinal symptoms may have been of autonomic origin. The combination of gastric dysmotility and autonomic disturbance has been shown to be correlated in a number of disorders such as primary gastric dysautonomia or irritable bowel syndrome (21,22). As well, constipation in multiple sclerosis may be related to autonomic dysfunction (23).

REFERENCES

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CONCLUSIONS

We believe our case represents an unusual manifestation of multiple sclerosis. We base this conclusion on the definitive diagnosis of multiple sclerosis, the absence of specific gastrointestinal disease and the therapeutic response of the nausea and vomiting to primary central acting promethazine hydrochloride and corticosteroids. Furthermore, the absence of any gastroenterological disease, such as inflammatory bowel disease, after more than five years of follow-up also supports our impression. Cases such as these should alert non-neurology physicians that persistent upper and lower gastrointestinal symptoms may, on occasion, be a consequence of primary central nervous system pathology.