foot and toes. During an episode, which lasted approximately 30 seconds, he complained of sharp pain in the left extremities that could not be explained solely by the muscle contraction causing the dystonic posturing. There were no alteration of consciousness, loss of sphincter control, or clonic movements in the limbs. The attacks had started without obvious precipitating factors 10 days before admission and occurred with a frequency of 5–10 per day.

The patient reported that since the age of 60, she had experienced a single similar attack of "painful spasm" in her left extremities that had not been treated and had not recurred.

Mild depressive symptoms had been present since the age of 60, being treated with fluoxetine and benzodiazepines intermittently. There was no family history of epilepsy or psychiatric disease. Her interictal general and neurological examination was unremarkable. Blood counts and serum biochemistry were normal. The routine EEG recording was normal. After administration of the awake EEG, disclosed no abnormalities; during drowsiness and sleep an almost continuous focal epileptiform activity was apparent in the right frontal-frontal-sagittal region (figure). Computerized tomography with and without contrast and MRI of the brain were performed and showed mild symmetric brain atrophy without any evidence of focal structural or metabolic changes.

Correlation of clinical features with the EEG allowed the exclusion of a variety of differential diagnoses. The patient was classified as having an apparent idiopathic epilepsy and a probable benign familial neonatal seizures.

The patient, because of the frequency of seizures, was advised to take a benzodiazepine; however, she refused treatment. After 15 years, she had only experienced one attack, and at age 75 she had further attacks. She was reassured that there was no evidence of progression of her condition, and the risks and benefits of treatment were discussed. The patient was advised to return if she experienced any further signs of seizures. She has been lost to follow-up since that time.

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Treatment of spasmodic torticollis with intramuscular phenol injection

Phenol is a caustic agent that produces tissue destruction and has been used to weaken muscle in patients with spasticity by either nerve block, motor point block, or intramuscular neurolysis.1,2 Local muscle pain and tenderness but no systemic or long term side effects have been reported after such use.3 Patient 1 was the first to use the intramuscular phenol in the treatment of spasmodic torticollis. I have used this agent in patients with spasmodic torticollis who have not responded well to intramuscular injections of botulinum toxin type A (BTX). Both patients gave informed consent to participate in the trial with the approval of the Duke University Institutional Review Board.

The muscles responsible for abnormal head movements were determined by clinical examination and by EMG recordings made with a concentric EMG needle in the sternomastoid, splenius capitis, and other neck muscles involved in head turning. Phenol solution (1% weight/volume USP phenol crystals in sterile aequous solution) was injected into the involved neck muscles with a recording monopolar injection electrode to determine that injections were made in muscle active in the abnormal movement. Injections were made at two to six sites in each muscle, depending on the motor unit action potentials with sharply rising components were recorded, indicating that the needle tip was close to muscle fibres that were activated during the abnormal movement. Areas near major vessels or nerves were avoided. No side effects or serious sequel injections were given if improvement was incomplete as determined by the patient's symptoms and by examination. At each visit, examination of strength in the involved muscles and functional assessment of the patient's torticollis were recorded on videotape for comparison with examinations made before phenol treatment.

Patient 1 is a 53 year old man who has had torticollis since age 43. Medical treatment had produced no improvement. At the age of 48, he had shown moderate improvement with BTX injections but, after 3 months, the responses disappeared. Injections diminished despite increasing doses. Muscles no longer developed weakness, atrophy, or denervation changes on EMG testing after BTX was injected. Antibody to botulinum could not be detected in the serum. He underwent selective peripheral denervation surgery when aged 50 and 51, with only mild improvement. Several months later, BTX injections were repeated with similar benefit.

He had tonic, uncontrollable turning of the head to the left with a phasic component and mild retrocollis before phenol injection. He maintained his head in the midline with great difficulty while sitting. Standing and any attempt to talk, walk, or use his hands produced immediate uncontrollable head turning. He had constant pain over the neck and intrascapular pain.

Within 18 hours after the initial injection of 100 mg phenol into the left splenius capitis, splenius cervicis, and longissimus capitis muscles, he noted a fuller range of motion in the neck and mild improved head control while walking. The injected muscles were tender and oedematous but the neck and intrascapular pain was considerably less. After subsequent injections he noted progressive improvement. After receiving a total dose of 500 mg of phenol over one month, he could walk, drive, or sit to eat in public with only occasional involuntary head turning. Maneouvres that previously had exacerbated the abnormal head movements now were performed with little or no difficulty. He estimated that pain was reduced by 90%. His only remaining problem is mild dystonia and tenderness over the injected muscles lasting one to two days after each injection. The improvement was sustained for five months after the initial injection. He then experienced gradual worsening of head control and some return of pain. He has subsequently received phenol injections at intervals of six months to maintain improvement.

Patient 2 is a 43 year old white male physician who had onset of involuntary head turning to the left at the age of 33. Medical treatment produced intolerable side effects and no significant improvement. At the age of 41, he received BTX injections every three to four months with moderate improvement but symptoms of head turning and pain returned four to six weeks after each injection. After each BTX injection, he developed severe dysphagia that persisted for two weeks. Before phenol was injected, he had moderately severe torticollis that was greatly improved by public speaking and manual activities.

Eighteen hours after the initial injection of 120 mg phenol into the left splenius capitis, splenius cervicis, and longissimus capitis muscles, he noted improvement in neck
pain but little change in head control. After subsequent injections of 260 mg given over two days, he noted considerable improvement in the ability to drive and carry out other manual tasks in his work. He graded improvement in head control and pain as 50% and 80%, respectively, above baseline, compared with 50% and 50% after the previous BTX injections. He had transient mild tenderness in the injected muscles. After the first month he experienced a slight decline (10%) in function and pain control that remained constant for the next four months and then gradually declined again. With subsequent injections of phenol at about six monthly intervals, he has maintained his maximal level of improvement.

These two patients had moderately severe spasmic torticollis that had improved only partially after previous treatment. In the first patient, BTX initially provided relief but became ineffective. In the second patient, BTX provided improvement but the side effect of dysphagia was nearly intolerable. Within 18 hours after phenol injections into cervical muscles, there was definite reduction of involuntary movements and pain, with functional improvement. Improvement was greater than after all previous treatments and persisted for six and five months respectively, after the initial series of phenol injections. The only side effect was transitory—namely, mild tenderness in the injected muscles.

In patients who become resistant to repeated injections of BTX, presumably due to formation of antibody to the toxin, it would be of great benefit to have an alternative treatment. Phenol may be of benefit in this situation and has the additional advantage of being inexpensive. If the promising response in these two patients is confirmed in a larger series of patients I am currently studying, EMG guided intramuscular phenol injections may prove to be an effective treatment for some patients with spasmic torticollis.

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Tuberculous myelopathy: a serial MRI study

In two definitive publications in 1969 Wadia and Dastur delineated spinal menigitides with associated radiculomyelopathy with particular reference to tuberculosis. The advent of MRI has meant that the nature of the intramedullary lesions can for the first time be defined during life. Furthermore, the natural history of spinal cord involvement in tuberculosis can be studied with serial MRI. The current study involves a single case followed up with a series of MRI examinations over an eight month period. It shows a previously unpublished pattern of cord involvement.

A 26 year old Samalian male refugee who had been in the United Kingdom for one year presented with a three month history of nausea and vomiting associated with increasingly severe headaches. He had no previous history of or exposure to tuberculosis. On admission to hospital he had neck stiffness only. Initial CSF examination
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